UST # 12458 - CAP

South Carolina Underground Storage Tank Management Division

Title: Programmatic QAPP Revision Number: 3.1 Revision Date: February 2016

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Appendix K. Contractor Checklist

For each report submitted to the UST Management Division, the contractor will be required to verify that all data elements for the required scope of work have been provided. For items not required for the scope of work, the N/A box should be checked. For items required and not completed or provided, the No box should be checked and a thorough description of the reason must be provided.

Item #	Item	Yes	No	N/A
1	Is Facility Name, Permit #, and address provided?	V		
2	Is UST Owner/Operator name, address, & phone number provided?	Vno	phone	# Ovo
3	Is name, address, & phone number of current property owner provided?	V	Francisc	710
4	Is the DHEC Certified UST Site Rehabilitation Contractor's Name, Address, telephone number, and certification number provided?	/		
5	Is the name, address, telephone number, and certification number of the well driller that installed borings/monitoring wells provided?			1
6	Is the name, address, telephone number, and certification number of the certified laboratory(ies) performing analytical analyses provided?			/
7	Has the facility history been summarized?	V		
8	Has the regional geology and hydrogeology been described?	V		_
9	Are the receptor survey results provided as required?			V
10	Has current use of the site and adjacent land been described?	V		
11	Has the site-specific geology and hydrogeology been described?	V		
12	Has the primary soil type been described?	V		
13	Have field screening results been described?			V
14	Has a description of the soil sample collection and preservation been detailed?			V
15	Has the field screening methodology and procedure been detailed?			/
16	Has the monitoring well installation and development dates been provided?			/
17	Has the method of well development been detailed?			
18	Has justification been provided for the locations of the monitoring wells?			V
19	Have the monitoring wells been labeled in accordance with the UST QAPP guidelines?		0	/
20	Has the groundwater sampling methodology been detailed?			/
21	Have the groundwater sampling dates and groundwater measurements been provided?	V	historic	data
22	Has the purging methodology been detailed?			/
23	Has the volume of water purged from each well been provided along with measurements to verify that purging is complete?			/
24	If free-product is present, has the thickness been provided?	V	historic	data
25	Does the report include a brief discussion of the assessment done and the results?		molecula	V
26	Does the report include a brief discussion of the aquifer evaluation and results?			/
27	Does the report include a brief discussion of the fate & transport models used?			/

South Carolina Underground Storage Tank Management Division

Title: Programmatic QAPP Revision Number: 3.1 Revision Date: February 2016

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Item #	Item	Yes	No	N/A
28	Are the site-conceptual model tables included? (Tier 1 Risk Evaluation)			
29	Have the exposure pathways been analyzed? (Tier 2 Risk Evaluation)			/
30	Have the SSTLs for each compound and pathway been calculated? (Tier 2 Risk Evaluation)			/
31	Have recommendations for further action been provided and explained?	V		
32	Has the soil analytical data for the site been provided in tabular format? (Table 1)		/	
33	Has the potentiometric data for the site been provided in tabular format? (Table 2)	V	nistori	c data
34	Has the current and historical laboratory data been provided in tabular format?	V	nistori	c date
35	Have the aquifer characteristics been provided and summarized on the appropriate form?			/
36	Have the Site conceptual model tables been included? (Tier 1 Risk Evaluation)			V
37	Has the topographic map been provided with all required elements? (Figure 1)	~		
38	Has the site base map been provided with all required elements? (Figure 2)	/		
39	Have the CoC site maps been provided? (Figure 3 & Figure 4)	V	GW on	M
40	Has the site potentiometric map been provided? (Figure 5)	/	historia	
41	Have the geologic cross-sections been provided? (Figure 6)	V	histori	
42	Have maps showing the predicted migration of the CoCs through time been provided? (Tier 2 Risk Evaluation)			V
43	Has the site survey been provided and include all necessary elements? (Appendix A)			V
44	Have the sampling logs, chain of custody forms, and the analytical data package been included with all required elements? (Appendix B)			V
45	Is the laboratory performing the analyses properly certified?			/
46	Has the tax map been included with all necessary elements? (Appendix C)			V
47	Have the soil boring/field screening logs been provided? (Appendix D)			V
48	Have the well completion logs, DHEC Form 2099, and DHEC Form 1903 been provided? (Appendix E)			~
49	Have the aquifer evaluation forms, data, graphs, equations, etc. been provided? (Appendix F)			V
50	Have the disposal manifests been provided? (Appendix G)			V
51	Has a copy of the local zoning regulations been provided? (Appendix H)			
52	Has all fate and transport modeling been provided? (Appendix I)			
53	Have copies of all access agreements obtained by the contractor been provided? (Appendix J)			/
54	Has a copy of this form been attached to the final report and are explanations for any missing or incomplete data been provided?	/		

UST # 12458- CAP

South Carolina Underground Storage Tank Management Division Title: Programmatic QAPP Revision Number: 3.1 Revision Date: February 2016 Page: 201 of 215

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CORRECTIVE ACTION PLAN

for

81 FOOD MART, UST PERMIT #12458 1502 EAST GREENVILLE STREET ANDERSON, ANDERSON COUNTY, SOUTH CAROLINA

Prepared for:

SOUTH CAROLINA DEPARTMENT OF HEALTH AND ENVIRONMENTAL CONTROL

Prepared by:

ENVIRORISK CONSULTANTS, INC.

UST SITE REHABILITATION CONTRACTOR #428



Post Office Box 945 Grayson, GA 30017 678-635-7360

Issue Date: September 7, 2017

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APPENDICES:

TABLES

Table 1 – Soil Analytical Data - not applicable

Table 2/3 – Potentiometric and Laboratory Data (historical)

Table 4 – Site Specific Target Levels

Table 5 – Timeline (text Section 4.6)

FIGURES

Figure 1 – Topographic Map

Figure 2 – Site Base Map

Figure 3 – Soil CoC Map- not applicable

Figures 4A-4D – Groundwater CoC Map (November 2016 data)

Figure 5 – Site Potentiometric Map (June 20, 2016 data)

Figures 6A-6C – Geologic Cross Sections (previous consultant)

Figure 7 – Proposed Treatment Plan



APPENDIX A: Site Survey- not applicable

APPENDIX B: Sampling Logs, Laboratory Data- not applicable

APPENDIX C: Tax Map- not applicable

APPENDIX D: Soil Boring/Field Screening Logs- not applicable

APPENDIX E: Well Completion Logs, Well Records- not applicable

APPENDIX F: Aguifer Evaluation Forms- not applicable

APPENDIX G: Disposal Manifests- not applicable

APPENDIX H: Local Zoning Regulations- *not applicable*

APPENDIX I: Fate and Transport Modeling- *not applicable*

APPENDIX J: Access Agreements- not applicable

APPENDIX K: Contractor Checklist

UIC Permit Application

Site Specific Work Plan

CORRECTIVE ACTION PLAN

for

81 FOOD MART, UST PERMIT #12458
1502 EAST GREENVILLE STREET
ANDERSON, ANDERSON COUNTY, SOUTH CAROLINA

GEOLOGY CERTIFICATION

I certify that I am a qualified ground-water scientist who has received a baccalaureate or post-graduate degree in the natural sciences or engineering, and have sufficient training and experience in ground-water hydrology and related fields, as demonstrated by state registration and completion of accredited university courses, that enable me to make sound professional judgments regarding ground-water monitoring and contaminant fate and transport. I further certify that this report was prepared by myself or by a subordinate working under my direction.



Kenneth C. Summerour, P.G. #2114 Registered Professional Geologist 9-7-17_____ Date



1.0 INTRODUCTION

This Corrective Action Plan (CAP) is being submitted for 81 Food Mart, UST Permit #12458, located at 1119 Cherokee Ave, Gaffney, South Carolina and Fast & Fresh 3, UST Permit #09530, located at 1502 East Greenville Street, Anderson, Anderson County, South Carolina (hereafter referred to as "the site" or "the facility"). The preparation of this CAP was authorized in Purchase Order (PO) #4600577312 for the release reported on November 10, 1992. This scope of work is being conducted for the South Carolina Department of Health and Environmental Control (SCDHEC).

This section outlines the purpose and content organization of the CAP along with a brief description of the site and surrounding properties.

1.1 Purpose

The overall purpose of this CAP is to present a plan for corrective action in order to prevent further degradation of the aquifer by continued migration of petroleum constituents into areas not previously impacted. Envirorisk has developed a corrective action scope of work to treat the Area of Concern (AOC) which includes all wells with measurable free product and dissolved concentrations. Corrective action at this facility described under this PO includes removal of free product to <0.01 feet and reduction of chemicals of concentration (CoC) to Site Specific Target Levels (SSTLs) as shown on **Table 4**. Historical data including water levels and analytical results are provided in **Table 2/3**.

All tables, figures, and other supplemental material referenced in the text are provided in the labeled appendices. Some tables, diagrams, etc. are provided in the body of the text for ease of reference.

1.2 Property Owner

The responsible property for UST #12458 is Brent Puzak with Circle K, and the mailing address is 305 Gregson Drive, Cary, NC, 27511. The phone number was not provided. The current property owner is Labtech Diagnostics, LLC, and their mailing address is 1502 East Greenville Street, Anderson, SC 29621 (same as the site address). The phone number is 864-760-0039.

1.3 Description of Site and Surrounding Properties

The site is currently operating as Labtech Diagnostics located on the south side of East Greenville Street at the intersection of McLees Road in Anderson, South Carolina. The site previously operated as a gasoline station/convenience store and contained three gasoline underground storage tanks (USTs) which were reportedly removed in April 1991. The quantity and cause of the reported release (dated November 10, 1992) is unknown.



The majority of the property is paved with concrete or asphalt with some landscaping around the perimeter. The surrounding area includes a mix of commercial and residential properties, with Sonic Drive-In Restaurant located south of the site, a former dry cleaner to the north, and a residence to the east. The site location including local topography is depicted on **Figure 1**. Site and surrounding properties are better illustrated on a site base map provided as **Figure 2**.

1.4 Summary of Previous Environmental Activities

Envirorisk reviewed technical files available electronically for the facility to gain an understanding of the site history. Currently, a total of thirty one (-31-) monitoring wells are included as part of the sampling network (see **Table 2/3**). Well locations are depicted on **Figure 2**. Shallow wells were installed to depths of 13-25 feet below ground surface (ft-bgs) with 10-foot screen sections. Deeper wells (indicated by a "D" at the end of the well ID or a "DMW" designation) were installed to depths of 47-112 ft-bgs with 5-foot screen sections. All wells were installed prior to Envirorisk's site involvement. Based on a file review, previous corrective action at the site has included multiple extraction events.

Based on the most recent gauging event conducted on November 21-23, 2016 free product was detected as follows: MW-1 (2.01'), MW-5 (0.83'), MW-6 (0.29'), MW-8 (0.29'), MW-9 (1.09'), MW-11 (0.35'), MW-12 (0.33'), MW-17 (0.07'), MW-23 (0.23'), and MW-30 (1.82'). Free product was only detected in the shallow wells. Groundwater samples were most recently collected in November 2016 for analysis of BTEX, naphthalene, MTBE, TAA, and TBA. The highest total concentration reported was 36,250 micrograms per liter (μ g/L) in MW-13, located south of the site near Sonic. **Table 2/3** (provided in the Appendix file from SCDHEC) is provided in the appendix and contains the tabulated data for November 2016. Groundwater CoC maps were created by a previous consultant for the November 2016 sampling event and are included as **Figures 4A-4D** for reference.

It is Envirorisk's understanding that horizontal and vertical delineation of the site has been completed by previous consultants to the satisfaction of SCDHEC. No additional site delineation is proposed in this CAP.

2.0 PHYSICAL SETTING

The physical setting of the site and surrounding region is described in this section. Discussions of regional characteristics were derived from published sources. Site specific characteristics, particularly geological classifications and unit interpolation, were based on a review of readily available file information obtained from SCDHEC.

2.1 Topography and Groundwater Flow Direction

The site elevation is approximately 699' above mean sea level and site groundwater flow, based on a review of previous groundwater flow maps, is to the south/southwest. The closest receptor (according to the 2000 Tier I Assessment Report) is Cox Creek located approximately 700 feet southwest of the site. **Figure 1**, attached, shows site topography. A potentiometric surface map, prepared by a previous consultant using June 2016 data, is included as **Figure 5**.

2.2 Regional and Site Geology and Hydrogeology

The subject site is located in the Piedmont Physiographic province of South Carolina. This province is characterized by rolling hills and stream valleys. In place weathering has chemically altered the bedrock and resulted in the residual soil retaining the structural features of the parent rock material. Bedrock of the area consists of layers of granite gneiss, biotite gneiss, and biotite schist. A review of boring logs in SCDHEC files indicates soils generally consist of clays, clayey silts, sandy silts, and fine to medium sands followed by partially weathered rock (sandy silt/rock fragments/micaceous gneiss bedrock). Geologic cross sections provided in SCDHEC records are included as **Figures 6A-6C**.

Shallow aquifers in the area usually occur in the saprolite unit which is hydraulically connected to the bedrock aquifer. Previous receptor surveys indicated no water supply wells within 1,000 feet of the site. Cox Creek was identified approximately 700 feet southwest and down-gradient of the site. During the November 2016 sampling event, groundwater in the surficial water bearing zone was encountered at depths ranging from approximately 6-14 ft-bgs.

2.3 Hydraulic Flow Characteristics

Hydraulic flow properties including hydraulic conductivity, hydraulic flow, and linear groundwater flow velocity or seepage velocity were evaluated by prior consultants. The hydraulic conductivity can be loosely defined as the velocity at which groundwater moves through the water-bearing soil medium.



The average hydraulic conductivity (K) of the surficial water bearing unit was previously calculated using the MW-2 and MW-3 at 17.0×10^{-5} and 28.29×10^{-5} feet per minute (ft/min), respectively. A seepage velocity of 2.86 feet per year was calculated based on an average K value of 22.65×10^{-5} ft/min (or 0.326 feet per day), a hydraulic gradient of 0.0084 feet per foot (ft/ft), and a porosity of 35% (sandy clay loam). The hydraulic gradient was calculated at 0.055 ft/ft in a July 1, 2014 Groundwater Monitoring Report.

3.0 CORRECTIVE ACTION SCOPE AND TECHNOLOGIES

The corrective action strategy developed for this site includes reduction of free product to <0.01' and reduction of dissolved constituents to the SSTLs as shown on **Table 4**. Prior to selecting a corrective action strategy, several technologies were evaluated based on Envirorisk's experience and a review of reported site conditions, as described in the subsections to follow. Based on the most recent data collected in November 2016, free product is present in ten monitoring wells in thicknesses ranging from 0.07 to 2.01 feet.

3.1 Summary of Remedial Site Conditions

The treatment area is shown on **Figure 7** and measures approximately 85' by 135' or 11,475 square feet and is located underneath the parking lot and a portion of the adjacent Right-of-Ways (ROWs) for East Greenville Street and McLees Road. The vertical extent, based on historic water table fluctuations, ranges from 6 to 14 ft-bgs in a fine to medium-grained silty-sand/sandy-silt saprolite. Based on a review of November 2016 data, monitoring wells located at or near the up-gradient plume boundary include MW-28, MW-7, and MW-27 to the north-northeast and MW-4, MW-23, and MW-25 to the south-southeast and down-gradient. Free product is present throughout the treatment area. Treatment areas are discussed in further detail in Section 4 and are shown on **Figure 7**.

3.2 Evaluation of Removal Technologies

Envirorisk evaluated various technologies to remediate the site to the SSTLs. The remedial technologies evaluated included soil vacuum extraction (SVE), air sparging (AS), dual-phase or multi-phase extraction (MPE), surfactant injections/extractions, chemical oxidation, enhanced bioremediation, and soil blending/excavation. Technologies may be combined with one or more methods as part of a full site corrective action strategy. A discussion of each technology is provided in the italicized sections.

3.2.1 Soil Vacuum Extraction (SVE)

SVE works by vacuum stripping volatile compounds out of interstitial soil pores through air movement without the use of groundwater extraction. Any volatile compound that exists in the vapor phase under ambient temperatures can theoretically be removed by SVE. Vacuum induced air flow in the vadose zone simply serves to enhance the volatilization process. The success of SVE at a given site is dependent on a number of factors, most notably soil permeability, moisture content, depth to water, fuel type, and contaminant distribution in the vadose zone versus free product presence beneath the water table.

A typical SVE system consists of a 5-10 horsepower (or greater) skid-mounted vacuum blower with a moisture separation tank and suitable high level float controls. Depending on air treatment requirements, influent vapors may need to be treated using thermal or carbon based methods prior to atmospheric discharge. Although this method is generally considered cost effective, it does not effectively address the presence of adsorbed free product trapped below the water table as a result of seasonal fluctuations. For this reason, SVE tends to have the greatest effectiveness in remediating "newer" free product releases in areas where the water table fluctuation has been minimal.

This technology was not considered in bid preparation and will not likely be utilized for site treatment. However, SVE may be considered for site treatment in combination with another technology such as air sparging described below.

3.2.2 Air Sparging

Air sparging consists of forcing compressed air below the water table in order to create transient air filled regimes in the saturated zone. Dissolved constituents exposed to the spared air environment are "stripped" from the dissolved phase into the gas phase where they can be captured as they migrate into the vadose zone with SVE technology. The air sparging process effectively creates a subsurface air stripper where the soil medium acts as the "packing" around which the injected air bubbles migrate through the water column. The sparging process also creates turbulence and increased subsurface mixing between soil and groundwater saturated zones, thereby resulting in the liberation of higher volatile organic compound (VOC) concentrations for SVE removal.

AS/SVE technology is generally effective for treatment of VOCs provided the compound has a Henry's Constant greater than 10⁻⁵. The success of AS/SVE at a given site is dependent on a number of factors most notably hydraulic conductivity, aquifer heterogeneity, soil permeability, depth to groundwater, ground cover, and contaminant distribution in the vadose zone and beneath the water table.

This technology may be utilized in conjunction with SVE as a follow-up technology.

3.2.3 Multi-Phase Extraction (MPE)

MPE is a technology that attempts to overcome the limitations of SVE, free product skimming, and standard pump-and-treat technologies through the simultaneous removal of volatile hydrocarbons from the dissolved phase, free product, vapor phase, and adsorbed soil particle phase. The term "multi-phase" extraction refers to the system's ability to extract free product, impacted groundwater, and soil vapors with specially designed high vacuum blowers capable of handling liquid and vapor streams. This method is similar in application to pump-and-treat but with the advantage of treating contaminated soils above and below the water table.

MPE is commonly applied through the use of vacuum trucks (AFVR) or a stationary system. Both applications involve drawing a high vacuum through PVC drop tubes installed in new or existing monitoring or recovery wells. The vacuum draws the water table down to an equilibrium level while extracting free product and adsorbed hydrocarbon constituents from the exposed soils beneath the water table. As the drop tubes are moved upward or downward, depending on the screened interval of the recovery wells, free product can be progressively removed from various portions of the soil column where impact is present including the vadose zone. The goal of the technology is to remove a minimal amount of groundwater while extracting free product and adsorbed hydrocarbons in the soil vapor. The success of the technology is determined by measuring the concentration of hydrocarbons in the off-gas stream and through measuring the resulting free product thicknesses and petroleum concentrations in the groundwater.

The success of MPE is dependent on soil permeability, moisture content, depth to water, fuel type, and contaminant distribution. A pilot test is generally performed prior to installing a stationary MPE system to determine the extent of groundwater drawdown and vacuum and flow rate needed for optimum recovery. A typical MPE system consists of a high vacuum liquid ring blower with a moisture separation tank and an oil-water separator and air stripper for influent groundwater. Depending on air treatment requirements, influent vapors may need to be treated using thermal desorption or similar method prior to atmospheric discharge.

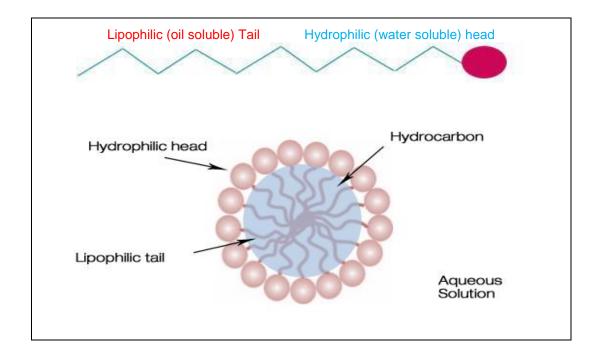
Mobile MPE will likely be utilized in combination with surfactant applications and in combination with other remedial methods.

3.2.4 Surfactant Injections/Extractions

Surfactants can be injected to reduce the surface tension between free product and soil particles where free product is typically "trapped" in interstitial pores. After reducing the surface tension, free product can be removed using simple groundwater extraction or mobile MPE (AFVR). Surfactants are chemical substances that have an affinity to both oil and water. The molecular structure of these compounds generally consists of a "Lipophilic" (oil/fuel soluble) tail and a "Hydrophilic" (water soluble) head as shown on the diagram on the following page.

When agitated, the two phases combine as microscopic oil/fuel droplets become encapsulated with a thin film of surfactant and water. These encapsulated droplets are referred to as micelles and when combined create stable macro-emulsions that break down the interfacial tension between trapped fuel/oil, pore water, and subsurface soil particles allowing more efficient recovery.

Based on the presence of free product and Envirorisk's prior successes using this remedial technology, surfactant treatments will be utilized for free product removal.

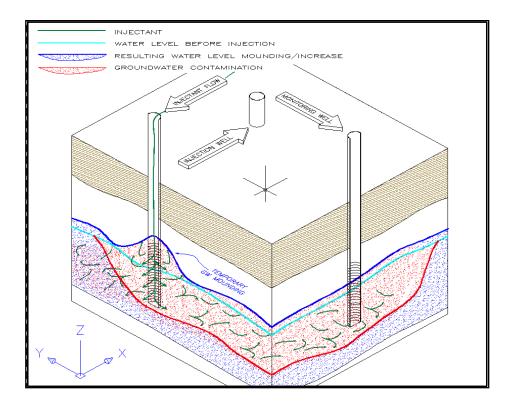


3.2.5 Chemical Oxidation

Chemical oxidation methods involve the use of concentrated oxidants to facilitate the chemical breakdown of hydrocarbons in the soil and groundwater. This chemical breakdown occurs as molecular bonds in organic compounds are "cleaved" and oxygen is inserted into the resulting fragments producing end products of carbon dioxide, water, and harmless salts. The oxidation process is generally driven by the creation of aggressive oxidant radicals that react on contact.

Chemical oxidants can be applied either in-situ via injection (ISCO), through in-place soil blending, or ex-situ by removing impacted soils for above ground treatment. Geological and hydrogeological conditions must be considered when developing the application strategy. ISCO field applications have clearly affirmed that matching the oxidant and delivery system to the contaminants of concern and the site conditions is the key to successful implementation.

ISCO field applications generally involve low pressure injection of oxidant fluid at various depths to provide treatment over the entire "smear zone" of contamination both above and below the water table. The addition of oxidant fluid produces a temporary "mounding" of the water table in the vicinity of the injection point. Since ISCO processes occur in the fluid phase, contaminants in the soil matrix must be desorbed from soils before they can react with the oxidant. A conceptual cross-section illustrating contaminant plumes targeted for injection is shown in the diagram below.



The primary drawback to the ISCO technology includes the need for complete oxidant contact with the contaminant which can be hampered by low permeability soils and preferential pathways in the soil matrix. Envirorisk relies on targeted injection techniques utilizing "positive placement monitoring" to overcome this drawback.

Contaminant removal can typically be facilitated at petroleum sites with the use of activated persulfate, catalyzed hydrogen peroxide (CHP), and/or use of solid phase peroxygen compounds including sodium percarbonate and calcium peroxide. Reactions for each are below.

Sodium persulfate oxidation occurs with and without activators including iron salts/chelates, an alkaline substance, induced heat, or hydrogen peroxide. The activation process results in the creation of sulfate radicals which have similar oxidant strength to hydroxyl radicals. The stylized oxidation reaction is as follows:

$$S_2O_8^{-2}$$
 + activator $\to SO_4^{-1}$ + $(SO_4^{-1} \text{ or } SO_4^{-2})$

CHP oxidation involves the combination of hydrogen peroxide, an iron catalyst, and a chelating agent resulting in the creation of hydroxyl radicals with a high oxidation potential.

The basic chemical reaction for CHP/Modified Fenton's oxidation is as follows:

$$H_2O_2 + Fe^{+2} + Chelating Agent \rightarrow Fe^{+3} + OH + OH + residual ionic salt$$

Modified Fenton's oxidation can also be produced using sodium percarbonate (Na₂CO₃•1.5H₂O₂). Sodium percarbonate is a soluble powder that releases hydrogen peroxide under mildly alkaline conditions (pH <9). The hydrogen peroxide generated can then be catalyzed with iron to produce hydroxyl radicals similar to a Modified Fenton's or CHP oxidant reaction. The chemical formula in solid phase and aqueous state for sodium percarbonate is shown below:

$$2Na_2CO_3.3H_2O_2 \rightarrow 2Na_2CO_{3(aq)} + 3H_2O_2$$

The hydrogen peroxide released will be react with either left over iron from previous ISCO treatments or supplemental iron added, resulting in Modified Fenton's oxidation:

$$H_2O_2 + Fe^{+2} \rightarrow Fe^{+3} + OH^- + OH.$$

The primary advantage of using sodium percarbonate to generate a CHP or Modified Fenton's reaction is to control the release of hydrogen peroxide and thereby extend the release of hydroxyl radicals. The oxidant reaction is non-corrosive and less exothermic than liquid based CHP applications. Following chemical oxidation, a release of dissolved oxygen will also occur to stimulate long term aerobic bioremediation.

Similar to sodium percarbonate, calcium peroxide can be utilized to generate a modified Fenton's reaction followed by aerobic bioremediation stimulated by a slow release of dissolved oxygen. Calcium peroxide has a low solubility and tends to form calcium hydroxide [Ca(OH)₂] at pH values of 11-12 as shown below:

$$2 CaO_2 + 2 H_2O \rightarrow Ca(OH)_2 + O_2$$

When the pH drops below 10 or 11, some hydrogen peroxide will be formed as shown in the reaction below:

$$CaO_2 + 2H + Ca_2 + (aq) + 2H_2O_2$$

Some or all of the hydrogen peroxide released will be used in a Modified Fenton's reaction as shown above with sodium percarbonate. Dissolved oxygen remaining after ISCO or from the dissolution of calcium peroxide under a higher pH will be subsequently utilized for aerobic bioremediation.

Based on site conditions and Envirorisk's prior success using ISCO for treatment of dissolved phase constituents, ISCO injections will be utilized for dissolved phase treatment.

3.2.6 Enhanced Bioremediation

Enhanced bioremediation is a process that attempts to accelerate the natural biodegradation processes already at work in the subsurface environment from microbial organisms. Enhancement is usually accomplished by providing nutrients, electron donors, or competent degrading microorganisms. The most common methods of enhanced bioremediation involve stimulating the subsurface by injecting nutrients to encourage either natural aerobic or anaerobic microbial activity. This is commonly referred to as in-situ bioremediation, or ISB.

Oxygen enhancement, commonly used to stimulate aerobic activity for degradation of petroleum hydrocarbons, can be achieved by either air sparging below the water table (bio-sparging including proprietary methods such as PHOSter™) or circulating low level oxygen enhancers throughout the contaminated ground water zone. Oxygen enhancement using bio-sparging is typically used in conjunction with SVE or bio-venting to enhance removal of volatiles. Additionally, solid-phase peroxide products or other slow release oxygenators can also be injected (ISB) into the subsurface to stimulate more rapid aerobic biodegradation.

Conditions influencing natural biodegradation need to be fully understood prior to enhancing natural processes. This is commonly performed by evaluating geochemical parameters in the groundwater along with relative concentrations of natural inorganics that may be utilized as electron acceptors and donors. Bench scale microbial studies may also be performed to better evaluate degradation rates.

Based on the contaminants present at this site (petroleum products), aerobic and/or anaerobic stimulation may be effective. Aerobic biostimulation will involve the use of oxygen release compounds in a "treatment train" approach. This can be accomplished using CHP, calcium peroxide, or sodium percarbonate oxidants that release oxygen during and following chemical oxidation. Anaerobic treatment would involve stimulating nitrate, iron, and/or sulfate-reducing bacteria in a reducing environment which may involve the injection of an organic carbon source to facilitate. Common organic substrates include sodium lactate or similar food grade products. Anaerobic treatment will be considered if sodium persulfate oxidation is utilized in order to take advantage of the residual sulfate remaining after the ISCO treatment.

Enhanced bioremediation, including ISB and bio-sparging technologies, may be utilized in combination with or following ISCO for site treatment of low to moderate dissolved phase constituents.

3.2.7 Soil Excavation/Soil Blending/Off-site Landfilling

In some cases, aggressive soil removal above and below the water table can be utilized as an effective means of removing impacted soils and blending in oxidants and/or bioremediation amendments. This process is often coupled with limited groundwater extraction to remove high dissolved phase constituents/free product recharging on the groundwater surface during excavating. Conventional excavation equipment has a maximum reach of 20 ft-bgs.

One alternative to conventional excavation is the advancement of a series of closely spaced large diameter borings using a "bucket" auger rig. This drill rig is similar to a caisson rig and is typically utilized in the installation of bored water wells. This drill method allows the removal of impacted soils with each advancement of the auger. Contaminated soils removed with the bucket attachment can be placed in roll-offs or dump trucks for off-site disposal at a Subtitle D landfill. In addition, impacted soils may be treated in-situ by blending in liquid and solid phase amendments to facilitate both ISCO processes in the groundwater and enhanced bioremediation. Temporary injection and/or extraction wells may also be installed in the open boreholes to further facilitate remediation.

Bucket auger rigs may be utilized in combination with ISCO treatment in the event that adequate contact and treatment is not achieved with ISCO and ISB.

4.0 PROPOSED PLAN FOR REMEDIATION

Based on the initial assessment of site conditions, free product and dissolved CoC reduction will be accomplished through a combination of surfactant injections/extractions and ISCO applications as outlined in the sections below. The primary injection technologies will be supplemented, if needed, with alternative technologies as described in Section 3. The treatment plan described below is shown on **Figure 7**. Per the Solicitation, physical access on the Sonic Restaurant is not included due to access issues. Therefore, SCDHEC has requested the use of angle-drilling beneath McLees Road ROW.

4.1 Drilling & Injection/Extraction Well Installation

To facilitate effective surfactant and ISCO product delivery, Envirorisk will advance a maximum of 25, two-inch diameter injection/extraction wells and a maximum of 55 temporary direct push injection points. The injection/extraction (IW/EWs) wells will be installed to depths of approximately 13 to 15 ft-bgs with 5-10 foot screens. Soil logging and field screening will be performed on a limited number of the IW/EWs to determine screen placement. The wells will be installed by a SC Licensed driller using either large diameter 3.25-inch inner diameter direct push rods or 2.25 to 3.25 inner diameter hollow stem augers. All wells will be constructed using Schedule 40 PVC with silica sand packs, hydrated bentonite seals, and secondary grout seals. The top of the PVC casings will be secured with a camlock fitting and threaded PVC cap to facilitate injection/extraction followed by a locking manhole cover enclosed in a concrete pad.

Following IW/EW installation and subsequent treatment, up to 55 direct push temporary injection points will be advanced for ISCO treatment. Depending on site conditions, some or all of these temporary injection points may be converted into one-inch diameter injection wells. The exact number of injection points/wells and locations will be based on field conditions, primarily including injectant flow and movement in the subsurface. Proposed IW/EW and direct push locations are displayed on **Figure 7**.

4.2 Surfactant Treatment/Extractions

A 12 to 24-hour mobile MPE/AFVR event will be initially performed using some or all of the newly installed injection/extraction wells. Extraction will be performed using one-inch drop tubes or "stingers" lowered to various depths below the water table in the contaminant "smear zone". The purpose of this initial mobile AFVR is to determine the quantity of fluid recovered from each IW/EW and/or group of wells for use in calculating surfactant dosing. In general, Envirorisk calculates surfactant doses assuming subsequent extraction removal of 1 to 1.5 times the liquid injected.

In a follow-up field mobilization, surfactant injection will be performed by injecting 200 to 500 gallons of a 2% to 5% solution into selected IW/EWs under mild pressure. After allowing sufficient time for free product emulsification and partial solubilization, one or more additional mobile AFVR events will be conducted to recover the emulsified free product and petroleum impacted groundwater. Waste manifests will be provided



following extraction events. Based on free product recharge and field conditions encountered, additional injection(s) and extraction(s) will be conducted to remove all or most of the free product.

During injection and extraction, Envirorisk will conduct field monitoring including depth to water and free product thickness gauging and geochemical parameter monitoring [pH, conductivity, temperature, dissolved oxygen (DO), and oxidation-reduction potential (ORP)], as needed. The degree of emulsification will also be evaluated through physical inspection of bailed free product and groundwater collected during treatment.

4.3 Angle-Drilled Wells

The Solicitation states that following free product removal, angle-drilled wells should be utilized for dissolved site treatment beneath McLees Road ROW. This requirement was presumably due to access restrictions associated with the Sonic Restaurant. A total of six or seven angled wells will be installed on the southeast portion of the treatment area as shown on **Figure 7.** Wells will be installed using 1" or 2" Schedule 40 PVC at an approximate 45-degree angle using a direct push rig or small auger rig. Screen sections will penetrate the subsurface within the impacted zone and will terminate in the midline of the road. Due to the angled installation, a silica sand pack will not be installed; however, a 2' prepack bentonite seal will be flush-threaded on top of the screen to prevent secondary grout intrusion. After grouting, wells will be completed with flush mounted well vaults. Wells will be utilized for ISCO treatment and/or mobile AFVR as described.

4.4 ISCO/Enhanced Bioremediation

Contaminant removal can typically be facilitated at petroleum sites with the use of activated persulfate, catalyzed hydrogen peroxide (CHP), and/or use of solid phase peroxygen compounds including sodium percarbonate and calcium peroxide. ISCO injections will be performed using up to 55 temporary injection points as well as up to 25 IW/EWs depicted on **Figure 7**. (The number of injection points/wells may be increased or decreased based on the effectiveness of the treatment). Based on experience with similar sites, the oxidant blend injected will likely consist of an initial treatment using CHP/Modified Fenton's (hydrogen peroxide and/or sodium percarbonate, calcium peroxide with an iron chelator) to reduce high dissolved VOCs and reduce/remove any remaining interstitial free product pockets. Following Modified Fenton's oxidation, unactivated or activated sodium persulfate will likely be injected to provide more long-term oxidant treatment. It is expected that two or more treatments will be required. Additional mobile AFVR events will also be performed, as needed, to supplement the ISCO treatments. The **UIC Permit** is included as an attachment.

Injection will be performed under low to moderate pressure using Envirorisk's mobile injection vehicle. During and immediately following ISCO injection, geochemical parameters (pH, conductivity, temperature, DO, and ORP) as well as depth to water and free product (if detected) thickness will be collected, as needed, to assess the radial extent of oxidant treatment.

Bioremediation (ISB) may be performed using solid or liquid phase amendments in conjunction with or following ISCO treatments. Amendments will be injected either with oxidants (i.e. same injection) or following. Alternatively, supplemental treatment may be performed using AS/SVE or bio-sparging (PHOSter) to reach lower dissolved constituent treatment goals. PHOSter is an air sparging technology in which triethylphosphate (TEP) is added through injections in the air stream to stimulate oxygen required to develop or sustain aerobic conditions. Implementation of these technologies may necessitate the need for additional air delivery injection wells as well as subsurface delivery piping. These systems are in-situ based and would NOT require off-site fluid discharges.

4.5 Semi-Annual Groundwater Sampling

As outlined in the Solicitation, all wells will be sampled semi-annually. If free product is detected the well will not be sampled. Wells will be sampled for the following constituents: BTEX, naphthalene, MTBE, TAA, and TBA using Method 8260. Sampling procedures will be followed as outlined in the "Quality Assurance Program Plan for the Underground Storage Tank Management Program, Revision 3.1" dated February 2016 and Envirorisk's Annual Contractor Quality Assurance Plan (ACQAP). As specified in the Solicitation, once free product has been removed, gauging will be conducted by SCDHEC. In addition, once sampling data indicates a 100% CoC concentration reduction, the Agency will be notified and corrective action activities will cease. The wells will be sampled for two quarters during the verification period. Split sampling will be performed with SCDHEC during the second verification event.

During the verification period, wells will be gauged and sampled quarterly. If the 100% CoC reduction goal has not been maintained, corrective action as described above will resume. Appropriate reports will be submitted following each gauging and/or sampling event.

4.6 Timeline

The table below lists predicted completion dates for the project. Dates are approximate, and are subject to change based on SCDHEC's input and response time to submittals, possible public notice delays, weather delays, holidays, changes in site conditions, and other conditions out of Envirorisk's control. The table is split into two sections, the first containing completed tasks and the second containing upcoming tasks.

Table 5. Timeline

	COMPLETED TASKS	
Task	Submittal	Date
Site Specific Work Plan for Initial Monitoring Event	Site Specific Work Plan	August 1, 2017
CAP Preparation / UIC Permit Application / Work Plan	CAP (this submittal)	September 7, 2017
	UPCOMING TASKS	
Task	Submittal	Date
SCDHEC review of CAP/ Public Notice	Notice to Proceed (from DHEC)	Assume 60 days- November, 2017
Initial Monitoring Event	Initial Monitoring Report	Due September 17, 2017
CAP Implementation	CAP Implementation Report	Field: 30 days from Notice to Proceed- December 2017 Report: 60 days from Notice to Proceed- January 2018 Invoice 40% of contract
Corrective Action / Semi- Annual Sampling	Appropriate Corrective Action System Evaluation (CASE) Report	Semi-annually with first report due within 90 days of CAP Implementation Report
Invoicing	CASE Reports	Invoice will be submitted with CASE reports for 10% of contract for reduction of free product to <0.01'
Invoicing	CASE Reports	Invoices will be submitted with CASE report for 10% of contract for CoC reduction of 60%
Invoicing	CASE Reports	Invoices will be submitted with CASE reports for 5% of contract for CoC reductions of 90% and 100%
Update QAPP Contractor Addendum	QAPP	First quarter of each year
Abandon and/or Remove Assessment and Corrective Action Equipment Components	Appropriate CASE Report	Within 60 days from notice by the Agency Invoice 30% of contract

A Contractor's Checklist is provided as **Appendix K**. The **Site Specific Work Plan** is also included as an attachment.





Site-Specific Work Plan for Approved ACQAP Underground Storage Tank Management Division

To:			(S	CDHEC Project Manager)
From:			(Co	ontractor Project Manager)
Contractor:		UST Contra	actor Certification Number:	· · · · · · · · · · · · · · · · · · ·
			UST Permit #	
Facility Address:				
Responsible Party:				
Current Use of Property	:			
Scope of Work (Please				
• '	☐ Tier II		☐ Groundwater Sampling	□ GAC
		Well Installation	□ Other	
Analyses (Please chec	k all that annly	<u> </u>		
Groundwater/Surface W		,		
☐ BTEXNMDCA (8260)		□ Lead	□ BOD	☐ Methane
☐ Oxygenates (8260B)	•	☐ 8 RCRA Metals	☐ Nitrate	☐ Ethanol
□ EDB (8011)		□ TPH	☐ Sulfate	☐ Dissolved Iron
□ PAH (8270D)		□ pH		
Drinking Water Supply V	Vells:	Δ р		
☐ BTEXNMDCA (524.2		☐ Mecury (200.8 245.	.1 or 245.2) □ EDB (504	l 1)
☐ Oxygenates & Ethan	•	• ,	,	,
Soil:	01 (02002)		,	
☐ BTEXNM ☐ Lead	I D RC	RA Metals	☐ TPH-DRO (3550B/8015B)) ☐ Grain Size
□ PAH		& Grease (9071)	☐ TPH-GRO (5030B/8015B	
Air:		S. S. Sudes (SS. 1)	(00002/00102	
□ BTEXN				
Sample Collection (Fs	timate the num	nber of samples of each i	matrix that are expected to be o	collected.)
Soil		Water Supply We		Field Blank
Monitoring W		Surface Water	Duplicate	Trip Blank
				r ·
Field Screening Metho	dology			
Estimate number and to	tal completed o	depth for each point, and	include their proposed location	s on the attached map.
# of shallow points propo	osed:	Esti	imated Footage:	feet per point
# of deep points propose	ed:	Esti	imated Footage:	feet per point
Permanent Monitoring	Wolle			
_		looth for each well, and i	include their proposed locations	on the attached man
	•	•	···	•
			d Footage:d footage:	
Comments, if warranted			l Footage:	ieet pei poliit
Comments, ii wananteu				
				· · · · · · · · · · · · · · · · · · ·

UST Permit #:	acility Name:		
Implementation Schedule (Number of Field Work Start-Up:	Field	roval) Work Completion: opies Provided to Property Owners:	
Aquifer Characterization Pump Test: □ Slug Test: □ (Chec	one and provide explar	ation below for choice)	
Investigation Derived Waste Disposa			
		Water: nase Product:	Gallons
Drilling Fluids:	_ Gallons Free-P	nase Product:	Gallons
Additional Details For This Scope of For example, list wells to be sampled, vevent, etc.		paired, well pads/bolts/caps to replace, detain	Is of AFVR
Name of Laboratory: SCDHEC Certification Number: _	? (Yes/No) If no, if no, if	indicate laboratory information below.	
Other variations from ACQAP. Pl	ase describe below.		
Attachments 1. Attach a copy of the relevant po	rtion of the USGS topog	raphic map showing the site location.	
Prepare a site base map. This must include the following: North Arrow Location of property lines Location of buildings Previous soil sampling location Previous monitoring well location Proposed soil boring locations Assessment Component Cost	Proposed monitorin Legend with facility Streets or highways Location of all presents Location of all potents	name and address, UST permit number, and (indicate names and numbers) ent and former ASTs and USTs ntial receptors	·

Table 2/3. Potentiometric and Laboratory Data Former 81 Food Mart, UST Permit #12458

Monitoring Well	тос	GW Depth (ft)	GW Elevation (ft)	FP (ft)	В	Т	E	Х	Naphth	MTBE	TAA	ТВА
MW-1 screen 15-25' Nov 21-23, 2016	94.87	9.75	83.61	2.01								
MW-4 screen 5-15' Nov 21-23, 2016	91.77	9.99	81.78		<5	< 5	< 5	< 5	<5	<5	<100	<100
MW-5 screen 5-15' Nov 21-23, 2016	94.34	11.68	82.04	0.83								
MW-6 screen 8-18' Nov 21-23, 2016	95.34	12.68	82.44	0.29								
MW-7 screen 5-15' Nov 21-23, 2016	95.80	10.17	85.63		<5	<5	<5	<5	<5	<5	<100	<100
MW-8 screen 5-15' Nov 21-23, 2016	95.80	10.15	85.43	0.29								
MW-9 screen 7-17' Nov 21-23, 2016	94.36	10.80	82.74	1.09								
MW-11 screen 7-17' Nov 21-23, 2016	94.71	11.40	83.05	0.35								
MW-12 screen 7-17' Nov 21-23, 2016	92.18	10.76	81.17	0.33								
MW-13 screen 3-13' Nov 21-23, 2016	90.12	9.71	80.41		8800	9500	3100	10000	650	<200	2800	<4000
MW-14 screen 15-25' Nov 21-23, 2016	NA	14.28	NA		<5	<5	<5	<5	<5	<5	<100	<100

Table 2/3. Potentiometric and Laboratory Data Former 81 Food Mart, UST Permit #12458

Monitoring Well	тос	GW Depth (ft)	GW Elevation (ft)	FP (ft)	В	Т	E	X	Naphth	MTBE	TAA	ТВА
MW-15 screen 15-25' Nov 21-23, 2016	NA	12.35	NA		<5	<5	<5	<5	<5	<5	<100	<100
MW-16 screen 10-20' Nov 21-23, 2016	NA	11.10	NA		<5	<5	<5	<5	<5	<5	<100	<100
MW-17 screen 10-20' Nov 21-23, 2016	NA	11.01	NA	0.07								
MW-18D screen 75-80' Nov 21-23, 2016	NA	11.36	NA		< 5	<5	< 5	<5	<5	< 5	<100	<100
MW-19D screen 43-48' Nov 21-23, 2016	NA	13.00	NA		<5	<5	<5	<5	<5	<5	<100	<100
MW-20D screen 42-47' Nov 21-23, 2016	NA	11.74	NA		< 5	<5	<5	<5	<5	<5	<100	<100
MW-21D screen 42-47' Nov 21-23, 2016	NA	10.14	NA		< 5	<5	<5	<5	<5	<5	<100	<100
MW-22D screen 42-47' Nov 21-23, 2016	NA	9.25	NA		< 5	<5	<5	<5	<5	<5	<100	<100
MW-23 screen 8-18' Nov 21-23, 2016	90.79	6.01	84.61	0.23								
MW-24 screen 10-20' Nov 21-23, 2016	91.49	9.88	81.61		900	80	120	360	52	810	2800	630
MW-25 screen 10-20' Nov 21-23, 2016	NA	12.65	NA		66	10	<5	91	25	<5	<100	<100

Table 2/3. Potentiometric and Laboratory Data Former 81 Food Mart, UST Permit #12458

Monitoring Well	тос	GW Depth (ft)	GW Elevation (ft)	FP (ft)	В	Т	E	Х	Naphth	MTBE	TAA	ТВА
MW-26 screen 10-20' Nov 21-23, 2016	NA	9.30	NA		< 5	< 5	< 5	<5	<5	<5	<100	<100
MW-27 screen 11-21' Nov 21-23, 2016	NA	11.75	NA		1700	1500	200	1100	100	42	240	<400
MW-28 screen 10-20' Nov 21-23, 2016	NA	11.12	NA		<5	<5	<5	<5	<5	<5	<100	<100
MW-30 screen 10-20' Nov 21-23, 2016	NA	10.01	NA	1.82								
DMW-1 screen 73-78' Nov 21-23, 2016	94.71	4.54	90.17		<5	<5	<5	<5	<5	<5	<100	<100
DMW-2 screen 75-80' Nov 21-23, 2016	NA	9.52	NA		<5	29	<5	<5	<5	<5	<100	<100
DMW-3 screen 107-112' Nov 21-23, 2016	NA	13.36	NA		<5	<5	<5	<5	<5	<5	<100	<100
KMO-17 screen 5-15' Nov 21-23, 2016	87.79	9.43	78.36		<5	<5	23	14	13	<5	<100	<100
KMO-18 screen 5-15' Nov 21-23, 2016	85.93	7.69	78.24		<5	<5	<5	<5	<5	<5	<100	<100

Notes

All analytical results are in micrograms per liter (μ g/L).

November 2016 data was collected by a previous consultant and was obtained from the Appendix file.

Groundwater elevations for wells containing FP are calculated as follows: TOC - [DTW + (FP thickness x 0.75)]

Screened intervals and TOC taken from SC DHEC appendix files.

NA = not applicable, TOC not provided

NS = not sampled

J = estimated value

ft = feet below ground surface

Table 4. Sitc specific Target Levels

81 Food Mart, UST Permit #12458

Site-Specific Target Levels (SSTLs) in parts per billion (ppb)

Well	Benzene	Toluene	Ethylbenzene	Xylenes	Naph.	MTBE	TAA	TBA
MW-1	40	26540***	3700***	21680***	285	1681	10086	58835
MW-4	5**	5**	5**	5**	5**	5**	100**	100**
MW-5	47	26540***	3700***	21680***	352	2339	14034	81865
MW-6	52	26540***	3700***	21680***	391	2757	16542	96495
MW-7	5**	5**	5**	5**	5**	5**	100**	100**
MW-8	52	26540***	3700***	21680***	391	2757	16542	96495
MW-9	27	26540***	3700***	21680***	185	862	5172	30170
MW-11	43	26540***	3700***	21680***	317	1984	11904	69440
MW-12	30	26540***	3700***	21680***	206	1020	6120	35700
MW-14	5**	5**	5**	5**	5**	5**	100**	100**
MW-15	5**	5**	5**	5**	5**	5**	100**	100**
MW-16	5**	5**	5**	5**	5**	5**	100**	100**
MW-17	62	26540***	3700***	21680***	481	3825	22950	133875
MW-18D	5**	5**	5**	5**	5**	5**	100**	100**
MW-19D	5**	5**	5**	5**	5**	5**	100**	100**
MW-20D	5**	5**	5**	5**	5**	5**	100**	100**
MW-21D	5**	5**	5**	5**	5**	5**	100**	100**
MW-22D	5**	5**	5**	5**	5**	5**	100**	100**
MW-23	19	12928	3700***	21680***	119	437	2622	15295
MW-24	23	80*	120*	360*	52*	437	2622	630*
MW-25	52	10*	5**	91*	25*	5**	100**	100**
MW-26	5**	5**	5**	5**	5**	5**	100**	100**
MW-27	25	1500*	200*	1100*	100*	42*	240*	400**
MW-28	5**	5**	5**	5**	5**	5**	100**	100**
MW-30	19	12,928	3700***	21680***	119	437	2622	15295
DMW-1	5**	5**	5**	5**	5**	5**	100**	100**
DMW-2	5**	29*	5**	5**	5**	5**	100**	100**
DMW-3	5**	5**	5**	5**	5**	5**	100**	100**

^{*} Lab analysis is less than calculated SSTLs, therefore SSTL mass is set equal to laboratory mass

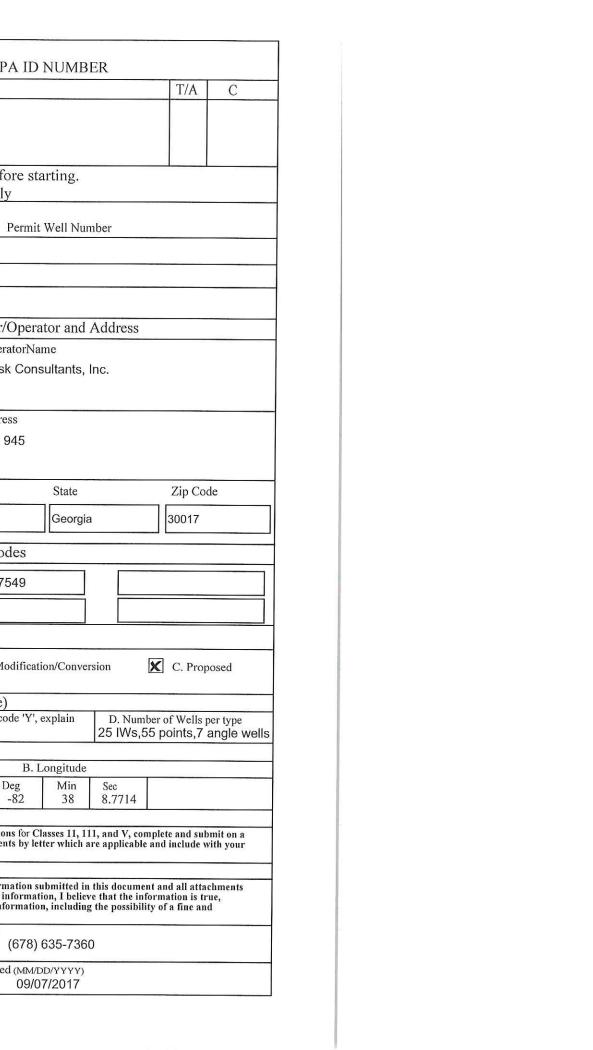
Physical corrective action on the Sonic Restaurant property is not included in this solicitation due to access issues. See Performance Requirement #2 for more details.

Source: SCDHEC Appendix file

^{**} Lab analysis is below detection limits, therefore SSTL mass is set equal to detection limits

^{***} SSTL is set to effective solubility limit

Form	DHE Ĉ				I. 1	EPA ID	NUMB	ER			_
Ĭ	Underground Injection Control							T/A	С		
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IV. Ownershi	p Status (Se	lect One)			V. SIC C	Codes					-
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D. Publi	с	E. Other (Explain)									Ī
VI. Well Statu											
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A. Opera	ating D	ate Started (MM	(/DD/YYYY)		B. Modification/Conversion C. Proposed						
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VII. Type of	Permit Red	quested - Class	and Type of	f Well	(see rever	se)		PT			_
A. Class(es) enter code(s) B. Type(s) enter code(s) C. If class is "oth					s code 'Y',	explain		per of Wells p			
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IX. Attachn	nents										_
Complete the follo separate sheet(s) a application.	owing question attachments A-	s on a separate sheet U as appropriate. At	(s) and number a tach maps wher	according e require	gly; see instruc d. List attachi	etions for Cl ments by let	asses 11, 11 ter which a	11, and V, con are applicable	plete and sub and include w	omit on a vith your	
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					B. Phone N		005 700	^			
Kenneth Summerour Principal Geologist				(678) 635-7360							
C Signature				D. Date Signed (MM/DD/YYYY)							



Instructions for Attachments to Form 1 Underground Injection Control for Corrective Action Wells (effective 01/91)

Attachment A: Activity for Review

Submit a brief description of the activities to be conducted that require a UIC permit.

Activities include drilling/installation of injection/extraction wells and surfactant injection(s) followed by extraction(s) and ISCO injection(s). Oxidants/chemicals/activators/bio-amendments utilized may include the following:

- sodium percarbonate;
- sodium persulfate;
- calcium peroxide;
- hydrogen peroxide; and
- sodium alpha olefin sulfonate.

MSDS forms are attached.

Attachment B: Well Construction Details

Submit schematics or other appropriate drawings of the surface and subsurface construction details of the recovery and injection wells.

Chemical will be applied via a maximum of 55 direct push injection points and/or a maximum of 25 temporary injection/extraction wells. In addition, up to 7 angled wells will be installed beneath McLees Road (per SCDHEC instructions in the Solicitation). Proposed locations are shown on Figure 7 and well schematics are attached.

Attachment C: Operating Data

Submit the following proposed operating data for each injection well:

1) Average and maximum daily rate and volume of fluid to be injected. In addition, indicate the average and maximum daily rate and volume of fluid to be withdrawn from each recovery well. Verification of the aquifer's hydraulic ability to product and accept the quantities proposed should be presented.

Injection quantities will vary from 100-500 gallons per injection well/point based on aquifer permeability. Typically, 2,000-3,000 gallons a day will be injected.

2) Average and maximum injection pressure.

10-50 psi.

3) Pumping schedule (i.e. continuous, alternating cycles, etc.)

N/A

UST #12458, Sept 2017 CAP

4) Proposed ranges in the concentration of all contaminant constituents within the injection fluid. Include comprehensive ground-water quality data from a "worst case" well sample.

Based on the most recent gauging event conducted on November 21-23, 2016 free product was detected as follows: MW-1 (2.01'), MW-5 (0.83'), MW-6 (0.29'), MW-8 (0.29'), MW-9 (1.09'), MW-11 (0.35'), MW-12 (0.33'), MW-17 (0.07'), MW-23 (0.23'), and MW-30 (1.82'). The highest total concentration reported was 36,250 micrograms per liter (µg/L) in MW-13, located off-site to the south. Site data is included on Table 2/3 in the CAP.

5) Length of time the project is expected to require injection to complete remediation (to ensure the effective dates of the permit will allow sufficient time to complete the project).

Approximately 2-5 years.

Attachment D: Monitoring Program

Discuss the planned monitoring program in detail:

1) Include a discussion of monitoring devices, sampling frequency (sufficient to verify treatment system efficiency), sampling protocol, sampling location, parameters to be analyzed, and proposed method(s) of analysis.

Wells will be sampled semi-annually. Wells containing free product will not be sampled. In addition, geochemical parameters (pH, conductivity, temperature, DO, and ORP) will be evaluated during injection activities.

2) This plan should indicate how, through monitoring, the proposed contaminant levels in the injectate will be verified.

Geochemical parameters will be collected during injection activities, and semi-annual groundwater sampling of all for analysis of BTEX, naphthalene, MTBE, TAA, and TBA.

3) This plan should also clearly illustrate exactly how hydraulic control of the contaminant plume (and injectate, where relevant) will be verified through monitoring (i.e., piezometers, quality analyses, etc.).

Perimeter wells will be sampled semi-annually to evaluate possible plume migration.

Attachment E: Existing of Pending State/Federal Permits

List the program and permit number of any existing State of Federal permits for the facility (i.e., NPDES, RCRA, UST, etc.).

UST Permit #12458

Attachment F: Description of Business

Give a brief description of the nature of the business of the facility and any immediately adjacent facilities.

UST #12458, Sept 2017 CAP

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UST #12458 is currently operating as Labtech Diagnostics, a medical diagnostics laboratory. The surrounding area includes a mix of commercial and residential properties, with Sonic Drive-In Restaurant located south of the site, a former dry cleaner to the north, and a residence to the east.

Attachment G: Area of Review

1) The area of review should be a fixed radius of ¼ mile from the injection well, the outermost injection wells (if a wellfield).

Your comment is noted.

2) If a fixed radius is not selected, the methods and the calculations used to determine the size of the area of review should be submitted.

N/A

Attachment H: Maps of Wells and Area of Review

1) Submit a topographic map of the area, extending one mile beyond the project boundaries. This map should show all hazardous waste treatment, storage, or disposal facilities, and all intake and discharge structures associated with the project facility. Any known areas of soil and/or ground-water contamination within a one mile radius should be indicated. Also indicate all surface water bodies of water, springs, mines (surface or subsurface), quarries, and other pertinent surface features such as residences, roads, and geologic faults (known or suspected).

Please refer to Figure 1. Site groundwater flow, based on a review of previous groundwater flow maps, is to the south/southwest. The nearest potential receptor is Cox Creek, located approximately 700' southwest of the site. No water supply wells were identified within 1,000 feet.

2) A scaled maps(s) should be included which shows the name and/or number and the location of ALL production, injection, monitoring, abandoned and dry wells within the area of review. This should be accomplished by file and field surveys. Information regarding the construction (i.e., total depth, diameter, casing/screened intervals, grouting, etc.) and the current status (i.e., actively used, temporarily abandoned, permanently abandoned) of ALL wells within the area of review should be submitted. If any wells have been abandoned, details on the method the wells were abandoned (i.e., cemented/grouted, filled with sand, etc.) should be included.

Please refer to Figure 2. A total of 31 monitoring wells are currently part of the sampling network at the site. Shallow well depths range from 13-25 feet with 10-foot screen sections and deeper well depths range from 47-112 feet with 5-foot screen sections. Additional construction details are available in SC DHEC files. Wells were installed prior to Envirorisk's involvement.

3) A potentiometric map of the project site should be submitted which accurately locates all monitoring wells and proposed recovery and injection wells and outlines the horizontal extent of both the free-phase contamination (where applicable) and dissolved contaminant plumes. Include all water level and product thickness data. The data and time that water levels and product thicknesses were measured should be indicated.

This map has not been prepared by Envirorisk for the CAP. However, a potentiometric map (Figure 5- June 2016 data) and plume maps (Figures 4A-4D- November 2016 data) were prepared by previous consultants and are attached. The most recent groundwater gauging and sampling event was conducted in November 2016.

Attachment I: Cross Sections/Diagrams

1) Geologic cross sections indicating the lithology and stratigraphy of the site and the horizontal and vertical extent of the contaminant plume, should be submitted. At least two stratigraphic cross sections, one parallel and one perpendicular to the horizontal ground-water flow direction, should be submitted.

This map has not been prepared by Envirorisk for the CAP. Previous cross sections prepared by other consultants are attached as Figures 6A-6C.

2) A schematic diagram, in the form of a cross section, showing the proposed remediation system with the components of flow (above and below ground) and all associated appurtenances (i.e., stripping tower, piping, wells, etc.).

N/A

Attachment J: Name and Depth of Underground Sources of Drinking Water (USDW's)

Identify and describe all aquifers which may be affected by the injection.

Envirorisk does not have detailed aquifer information; however, only groundwater in the immediate vicinity of the injection wells/points should be affected.

Attachment K: Hydraulic Control

1) Sufficient supporting data (i.e. time/drawdown data, Theis curves and methods, calculations, etc.), used to determine aquifer characteristics to verify complete hydraulic control over the contaminant plume (and injectate, if proposed injectate quality does not conform to classified ground-water standards) during injection should be submitted. At a minimum, values should be given for transmissivity, hydraulic conductivity, effective porosity and specific yield

Injection will not exceed pore volumes in the contaminated water zone. Injection will be performed from the perimeter into the heart of the plume to limit potential migration. The average hydraulic conductivity of the surficial water bearing unit (calculated by previous consultants) for MW-2 and MW-3 was calculated at 17.0 x 10⁻⁵ and 28.29 x 10⁻⁵ feet per minute (ft/min), respectively. The seepage velocity was previously reported at 2.86 feet/year. The hydraulic gradient was calculated at 0.055 ft/ft in a July 1, 2014 Groundwater Monitoring Report.

2) Demonstrate the presence and magnitude of, or the absence of, any vertical hydraulic gradient at the site. If a vertical hydraulic gradient exists, show how its direction and magnitude are incorporated in the calculations demonstrating hydraulic control.

Vertical gradients are not considered significant at this facility based on a review of file data.

UST #12458, Sept 2017 CAP

**Please note- per the Solicitation, migration of the dissolved plume has occurred to the south onto the Sonic Restaurant property. Sonic has denied physical access to the site; therefore, SCDHEC has requested the use of angled wells beneath McLees Road for treatment to the south of the site.

3) Ground-water flow computer models (especially 2-D may view with potentiometric and flow lines) may be utilized and submitted. All calculations should be in English units. All model-derived data and maps should be properly labeled and keyed so as to be clearly understood.

N/A

ATTACHMENT B: Well Construction Details

ENVIRORISK CONSULTANTS, INC.

INJECTION/EXTRACTION WELL CONSTRUCTION DETAILS (Example)

DRILLING FIRM:

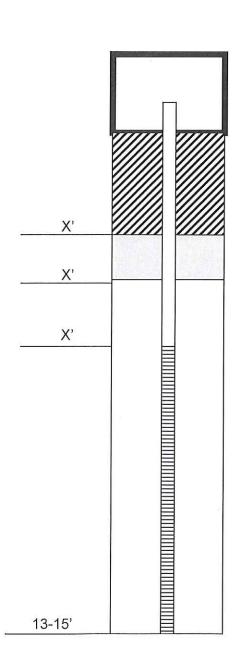
SUPERVISING GEOLOGIST:

PROJECT NAME:

PROJECT LOCATION:

INJECTION WELL NO.:

DATE OF INSTALLATION:



COMPLETED WITH STEEL, BOLT-DOWN WELL VAULT

WELL RISER

MATERIAL:

1 OR 2 INCH

DIAMETER: JOINT TYPE:

FLUSH THREADED

LENGTH:

X FEET

PVC

SECOND SEAL

TYPE OF SEAL: THICKNESS:

PORTLAND CEMENT X FEET

FIRST SEAL

TYPE OF SEAL:

BENTONITE CHIPS

THICKNESS:

2 FEET

FILTER PACK

TYPE OF FILTER:

SAND

DISTANCE ABOVE SCREEN:

2 FEET

WELL SCREEN

SCREEN MATERIAL:

PVC

DIAMETER:

1 or 2 INCH

LENGTH:

5-10 FEET

SLOT SIZE:

0.010 INCHES

DEPTH TO BOTTOM OF INJECTION WELL 13-15 FEET

ATTACHMENT B: Well Construction Details

ENVIRORISK CONSULTANTS, INC.

ANGLED INJECTION WELL CONSTRUCTION DETAILS (Example)

DRILLING FIRM:

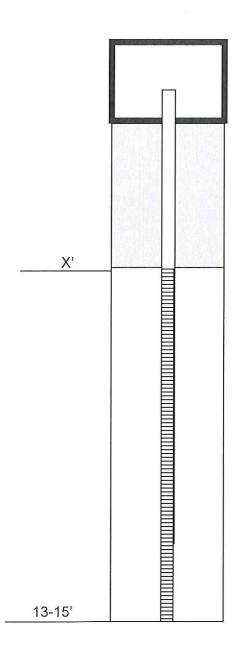
SUPERVISING GEOLOGIST:

PROJECT NAME:

PROJECT LOCATION:

INJECTION WELL NO.:

DATE OF INSTALLATION:



COMPLETED WITH STEEL, BOLT-DOWN WELL VAULT

WELL RISER

MATERIAL: DIAMETER:

PVC 1 OR 2 INCH

JOINT TYPE: LENGTH:

FLUSH THREADED 21 FEET

FIRST SEAL

TYPE OF SEAL: THICKNESS:

BENTONITE CHIPS

2 FEET

PRE-PACKED WELL SCREEN WITH SAND

SCREEN MATERIAL:

<u>PVC</u> 1 or 2 INCH

DIAMETER:

X FEET

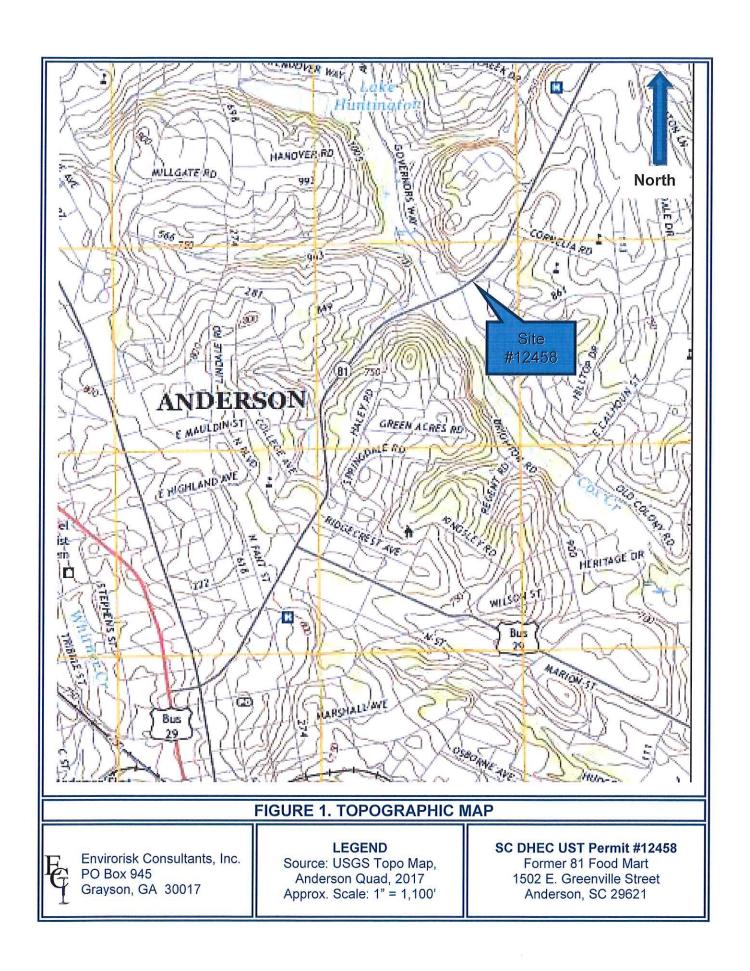
LENGTH: SLOT SIZE:

0.010 INCHES

DEPTH BELOW GROUND SURFACE

15 FEET

*Well will be installed at an approximate 45-degree angle equaling a 15-foot depth below ground surface using approximately 21-feet of well pipe.



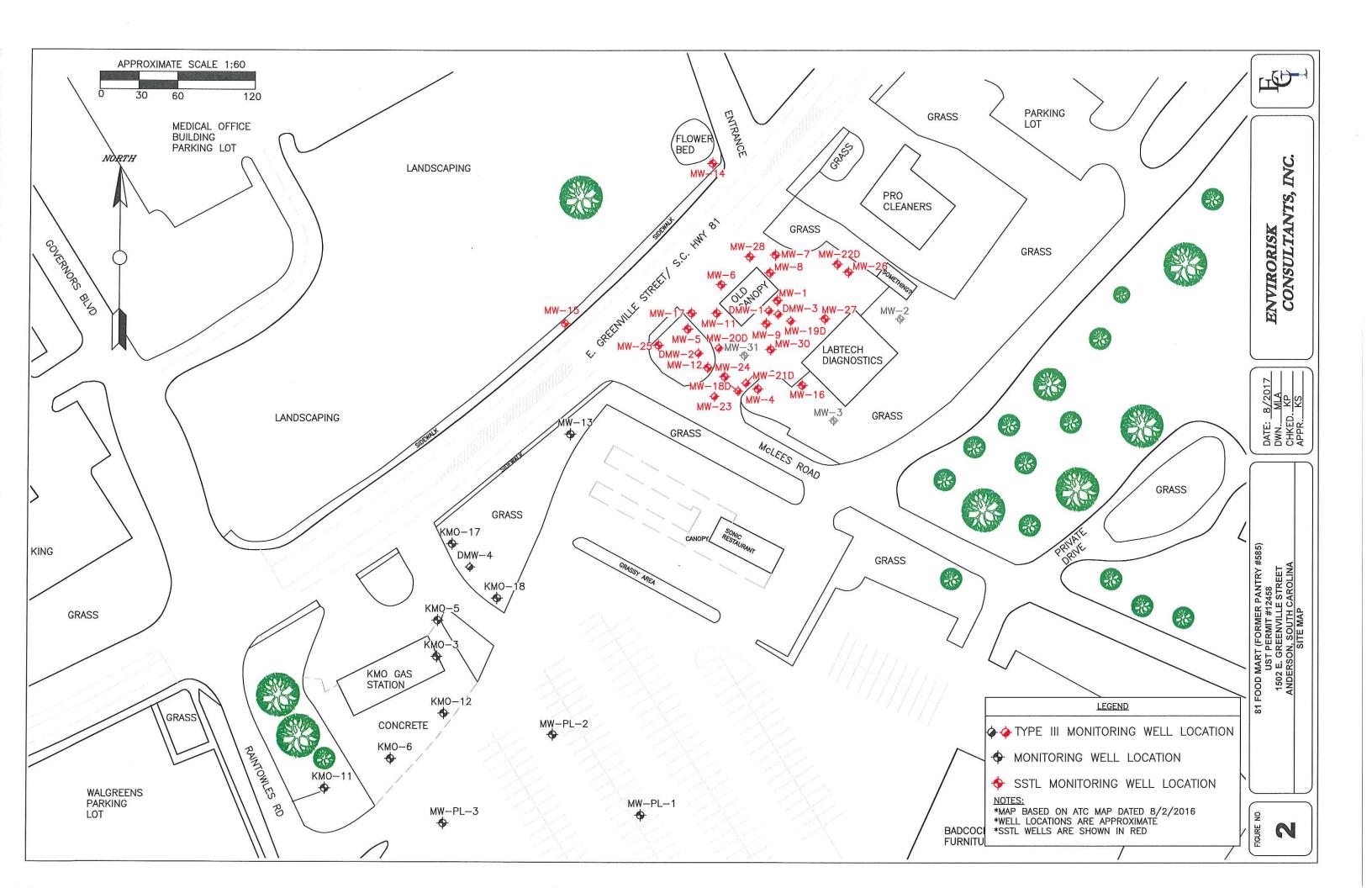
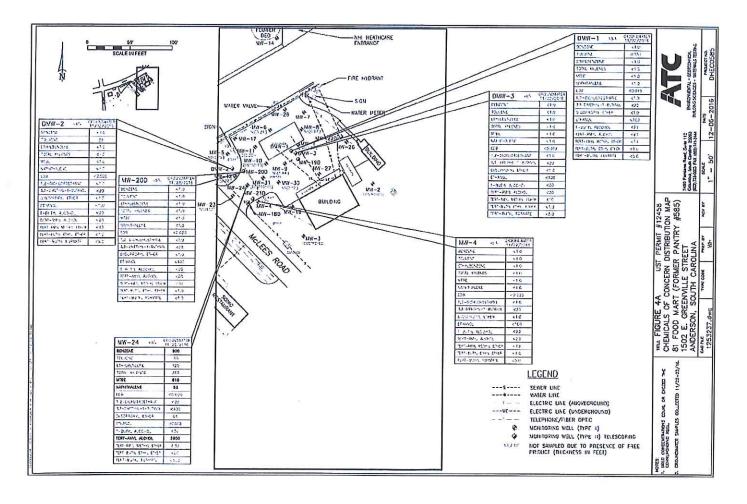


Figure 4A. Groundwater COC map, Nov. 2016



Source! SCOHEC technical file

Figure 4B. Groundwater Col map, Nov. 2016

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Source: SCDHEC technical file

| MINOSCHING| | MANUAL | MANUA

Figure 4C. Groundwater Coc map, Nov. 2016

Source: SCDHEC technical file

| Fig. |

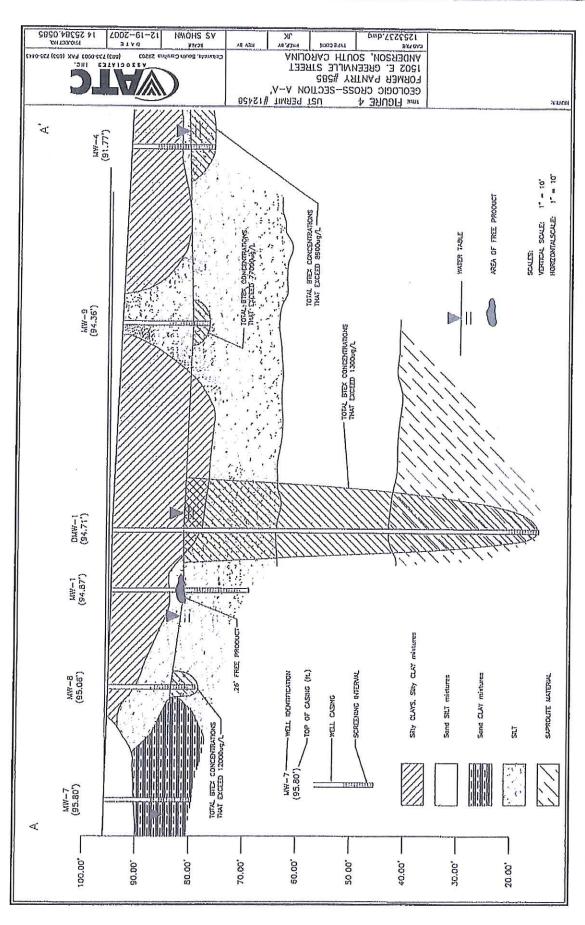
Figure 40. Groundwater Coc map. Nov. 2016

Source: SCDHEC technical file

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Higure					189. M.	© 1 - DRX VIOLETTA

Geologic Cross Section, Transect map Figure 64.

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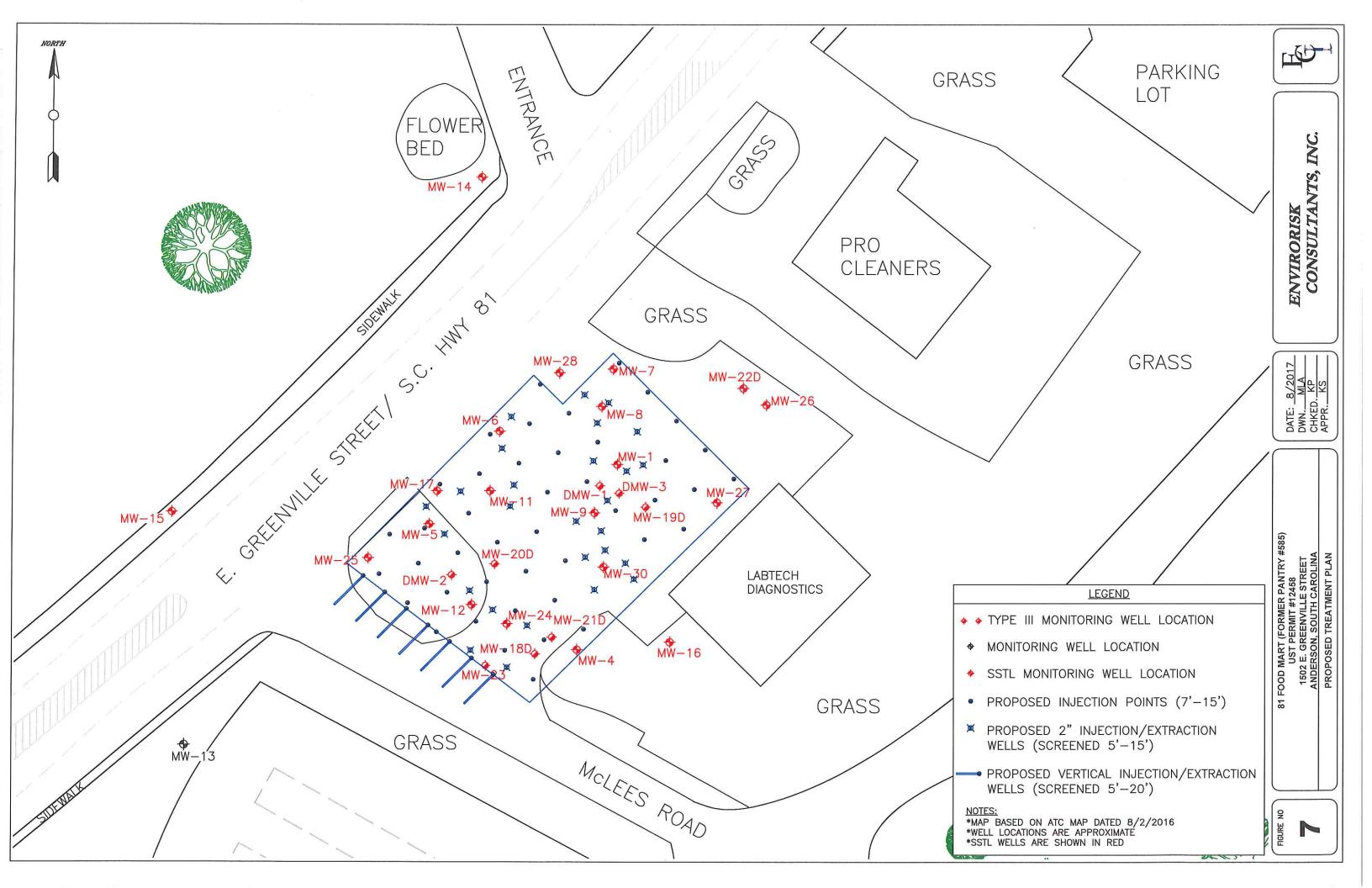


Sowre: SCDHEC Appendix file

13. Geologic Cross Section, A-A!
13. Geologic Cross Section,
13. Geologic Cross
B. Geologic
5.
9
Figure

Source: SCDHEC Appendix file

Figure Loc. Geologic Cross Section, 8-8'



HARCROS

MATERIAL SAFETY DATA SHEET C-13 A05-40

1. Product and Company Identification

Material name

T Det AOS-40

Version#

09

Revision date

05-08-2012

Chemical name

Chemical description

Sodium Alpha Olefin Sulfonate

CAS#

Organic Mixture

Mixture

MSDS Number

001133-09

Manufacturer information

Harcros Chemicals Inc 5200 Speaker Rd.

Kansas City, KS 66106 United States Main Telephone Number 1-913-321-3131

CHEMTREC -For Emergencies Only 1-703-527-3887 (call collect)

CHEMTREC -For Emergencies Only 1-800-424-9300

2. Hazards Identification

Emergency overview

Irritating to eyes.

OSHA regulatory status

This product is considered not hazardous under 29 CFR 1910.1200 (Hazard Communication).

Potential health effects

Routes of exposure

Eye contact.

Eyes

May cause eye irritation.

May be irritating to the skin.

Skin Inhalation

May cause irritation of respiratory tract.

Ingestion

May be harmful if swallowed. May cause irritation, coughing, headache, nausea, vomiting,

diarrhe

Potential environmental effects

May cause long-term adverse effects in the environment.

3. Composition / Information on Ingredients

Non-hazardous components	CAS#	Percent
Water	7732-18-5	56 - 64
SULFONIC ACIDS, C14-16-ALKANE HYDROXY AND C14-16-ALKENE, SODIUM SALTS	68439-57-6	36 - 42
Sodium Chloride	7647-14-5	0 - 1
Sodium Sulphate	7757-82-6	0 - 1

4. First Aid Measures

First aid procedures

Eye contact

Rinse immediately with plenty of water, also under the eyelids, for at least 15 minutes. Get

medical attention if symptoms occur.

Skin contact

Immediately flush with plenty of water for at least 15 minutes while removing contaminated

clothing and shoes. Wash clothing separately before reuse. Call a physician if systems develop or

persist.

Inhalation

If breathing is difficult, remove to fresh air and keep at rest in a position comfortable for breathing. Call a physician if symptoms develop or persist.

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MSDS US 1/6



Ingestion

Rinse mouth. Drink plenty of water. Get medical attention if symptoms occur.

General advice

If you feel unwell, seek medical advice (show the label where possible).

5. Fire Fighting Measures

Flammable properties

The product is not flammable. No unusual fire or explosion hazards noted.

Extinguishing media

Suitable extinguishing

media

Dry chemical, CO2, or water spray. Foam.

medic

Unsuitable extinguishing

media

Do not use water jet as an extinguisher, as this will spread the fire.

Fire fighting

equipment/instructions

Evacuate the area promptly. As in any fire, wear self-contained breathing apparatus pressure-demand, MSHA/NIOSH (approved or equivalent) and full protective gear. Cool containers exposed to heat with water spray and remove container, if no risk is involved.

Hazardous combustion products

Carbon Oxides. Sulfur Oxides.

6. Accidental Release Measures

Personal precautions

Keep unnecessary personnel away. Local authorities should be advised if significant spillages cannot be contained.

Environmental precautions

Prevent further leakage or spillage if safe to do so. Do not contaminate water.

Methods for containment

Stop the flow of material, if this is without risk. Dike the spilled material, where this is possible.

Methods for cleaning up

Should not be released into the environment.

Large Spills: Stop the flow of material, if this is without risk. Dike the spilled material, where this is possible. Absorb in vermiculite, dry sand or earth and place into containers. Following product recovery, flush area with water.

Small Spills: Wipe up with absorbent material (e.g. cloth, fleece). Clean surface thoroughly to remove residual contamination.

Never return spills in original containers for re-use. For waste disposal, see section 13 of the MSDS.

7. Handling and Storage

Handling

Avoid release to the environment. Do not breathe dust/fume/gas/mist/vapors/spray. Do not get in

eyes, on skin, on clothing.

Storage

Keep away from food, drink and animal feedingstuffs. Keep away from heat, sparks and open flame. Keep container tightly closed and dry. Keep containers between 50 F. and 100F. Use care in handling/storage. Store in accordance with local/regional/national/international regulation.

8. Exposure Controls / Personal Protection

Personal protective equipment

Eye / face protection

Avoid contact with eyes. Wear safety glasses with side shields (or goggles). Eye Wash Fountain

should be available.

Skin protection

considerations

Material name: T Det AOS-40

Wear suitable protective clothing and gloves.

Respiratory protection

No personal respiratory protective equipment normally required. If engineering controls do not maintain airborne concentrations below recommended exposure limits (where applicable) or to an acceptable level (in countries where exposure limits have not been established), an approved respirator must be worn.

General hygiene

Avoid contact with eyes. Handle in accordance with good industrial hygiene and safety practice.

9. Physical & Chemical Properties

Appearance

Yellow Amber

Physical state

Liquid.

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MSDS US 2/6

Liquid. Form Color Not available. Odor Slight. Odor threshold Not available. 7 - 9 25 mm Hg @ 77 F. Vapor pressure Vapor density Not available. **Boiling** point 212 °F (100 °C) Melting point/Freezing point Not available. Solubility (water) Soluble in all portions Specific gravity 1.05 Relative density Not available. Flash point Not available. Flammability limits in air, upper, Not available. % by volume Flammability limits in air, lower, Not available. % by volume Auto-ignition temperature Not available. Percent volatile 56 - 64 % 10. Chemical Stability & Reactivity Information Chemical stability Material is stable under normal conditions. Conditions to avoid Do not mix with other chemicals. Incompatible materials Strong oxidizing agents. Hazardous decomposition No hazardous decomposition products are known. products Possibility of hazardous Not established. reactions 11. Toxicological Information Toxicological data **Product** Species Test Results T Det AOS-40 (Mixture) Acute Dermal LD50 Rabbit 6300 mg/kg, CAS# 68439-57-6 Oral LD50 Rat 10000 mg/kg, Sodium Sulfate (Solution) 2310 mg/kg, CAS# 68439-57-6 Components Species **Test Results** Sodium Chloride (7647-14-5) Acute Oral LD50 Mouse 4000 mg/kg Rat 3000 mg/kg Other LD50 Mouse 2602 mg/kg Material name: T Det AOS-40 MSDS US 2691 Version #: 01 Revision date: 05-08-2012 Issue date: 05-08-2012

MSDS US 3 / 6

	Specie	S	Test Results
Sodium Sulphate (7757-82-	-6)		
Acute			
Other			
LD50	Rabbit		> 4 g/kg
* Estimates for product	may be based on	additional component data not shown.	
Local effects	May be ir	ritating to eyes.	
Carcinogenicity	This prod	uct is not considered to be a carcinogen by	IARC, ACGIH, NTP, or OSHA.
Further information	This prod	uct has no known adverse effect on human	health.
12. Ecological Information	tion		
Ecotoxicological data Product		Species	Test Results
T Det AOS-40 (Mixture)		Species	rest Results
Crustacea	EC50	Daphnia	10.6146 mg/l, 48 hours, estimated
Components	2000	Species	Test Results
Sodium Chloride (7647-14-5	5)	Openies	Test Results
Aquatic	2)		
Crustacea	EC50	Water flea (Daphnia magna)	340.7 - 469.2 mg/l, 48 hours
Fish	LC50	American eel (Anguilla rostrata)	0 - 27260 mg/l, 96 hours
Sodium Sulphate (7757-82-6		ransinan ser (rangama reenata)	5 El Essingia, so nodis
Aquatic	-,		
Crustacea	EC50	Water flea (Ceriodaphnia dubia)	2807 - 3535 mg/l, 48 hours
Fish	LC50	Striped bass (Morone saxatilis)	56 mg/l, 96 hours
SULFONIC ACIDS, C14-16		XY AND C14-16-ALKENE, SODIUM SALT	V250 0-
Aquatic		, ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	C (00 100 01 0)
			4.14 - 4.95 mg/l, 48 hours
Crustacea	EC50	Water flea (Ceriodaphnia dubia)	The most many to mound
* Estimates for product	may be based on	additional component data not shown,	•
	may be based on	additional component data not shown, uct complies with the biodegradability criter	•
* Estimates for product	may be based on This Prod 648/2004	additional component data not shown, uct complies with the biodegradability criter	•
* Estimates for product Ecotoxicity	may be based on This Prod 648/2004 An enviro	additional component data not shown, uct complies with the biodegradability criter nmental hazard cannot be excluded in the	ria as laid down in Regulation (EC) No.
* Estimates for product Ecotoxicity Environmental effects	may be based on This Prod 648/2004 An environ ty Not availa	additional component data not shown, uct complies with the biodegradability criter nmental hazard cannot be excluded in the	ria as laid down in Regulation (EC) No.
* Estimates for product Ecotoxicity Environmental effects Persistence and degradability	may be based on This Prod 648/2004 An environ ty Not availa Readily B	additional component data not shown, uct complies with the biodegradability criter nmental hazard cannot be excluded in the	ria as laid down in Regulation (EC) No.
* Estimates for product Ecotoxicity Environmental effects Persistence and degradability Chemical fate information	may be based on This Prod 648/2004 An enviror ty Not availa Readily B ations Collect an this mater or dispose Under RC disposal, v	additional component data not shown, uct complies with the biodegradability criter nmental hazard cannot be excluded in the cible. iodegradable, d reclaim or dispose in sealed containers a ial to drain into sewers/water supplies. This ad of, is not a hazardous waste according to RA, it is the responsibility of the user of the	ria as laid down in Regulation (EC) No. event of unprofessional handling or disposal. at licensed waste disposal site. Do not allow as product, in its present state, when discarded to Federal regulations (40 CFR 261.4 (b)(4)).
* Estimates for product Ecotoxicity Environmental effects Persistence and degradabilit Chemical fate information 13. Disposal Considera	may be based on This Prod 648/2004 An environ ty Not availa Readily B ations Collect an this mater or dispose Under RC disposal, with all ap	additional component data not shown, uct complies with the biodegradability criter nmental hazard cannot be excluded in the dible. iodegradable, directain or dispose in sealed containers a ial to drain into sewers/water supplies. This ad of, is not a hazardous waste according to RA, it is the responsibility of the user of the whether the product meets RCRA criteria for plicable regulations.	ria as laid down in Regulation (EC) No. event of unprofessional handling or disposal. at licensed waste disposal site. Do not allow as product, in its present state, when discarded to Federal regulations (40 CFR 261.4 (b)(4)). be product to determine, at the time of

 Material name: T Det AOS-40
 MSDS US

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14. Transport Information

Not regulated as dangerous goods.

IATA

Not regulated as dangerous goods.

IMDG

Not regulated as dangerous goods.

15. Regulatory Information

US federal regulations

This product is not known to be a "Hazardous Chemical" as defined by the OSHA Hazard

Communication Standard, 29 CFR 1910.1200.

All components are on the U.S. EPA TSCA Inventory List.

CERCLA/SARA Hazardous Substances - Not applicable.

Drug Enforcement Administration (DEA). List 2, Essential Chemicals (21 CFR 1310.02(b) and 1310.04(f)(2)

Not regulated.

DEA Essential Chemical Code Number

Not regulated.

Drug Enforcement Administration (DEA). List 1 & 2 Exempt Chemical Mixtures (21 CFR 1310.12(c))

Not regulated.

DEA Exempt Chemical Mixtures Code Number

Not regulated.

CERCLA (Superfund) reportable quantity

None

Superfund Amendments and Reauthorization Act of 1986 (SARA)

Hazard categories

Immediate Hazard - Yes

Delayed Hazard - No

Fire Hazard - No

Pressure Hazard - No

Reactivity Hazard - No

Section 302 extremely

hazardous substance

No

Section 311 hazardous

No

chemical Inventory status

Country(s) or region	Inventory name	On inventory (yes/no)*
Australia	Australian Inventory of Chemical Substances (AICS)	Yes
Canada	Domestic Substances List (DSL)	Yes
Canada	Non-Domestic Substances List (NDSL)	No
China	Inventory of Existing Chemical Substances in China (IECSC)	Yes
Europe	European Inventory of Existing Commercial Chemical Substances (EINECS)	Yes
Europe	European List of Notified Chemical Substances (ELINCS)	No
Japan	Inventory of Existing and New Chemical Substances (ENCS)	No
Korea	Existing Chemicals List (ECL)	Yes
New Zealand	New Zealand Inventory	Yes
Philippines	Philippine Inventory of Chemicals and Chemical Substances (PICCS)	Yes

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Country(s) or region

Inventory name

On inventory (yes/no)*

United States & Puerto Rico Toxic Substances Control Act (TSCA) Inventory

Yes

*A "Yes" indicates that all components of this product comply with the inventory requirements administered by the governing country(s)

State regulations

California Safe Drinking Water and Toxic Enforcement Act of 1986 (Proposition 65): This material is not known to contain any chemicals currently listed as carcinogens or reproductive toxins.

US - Pennsylvania RTK - Hazardous Substances: Listed substance

Sodium Sulphate (CAS 7757-82-6)

16. Other Information

Further information

HMIS® is a registered trade and service mark of the NPCA.

HMIS® ratings

Health: 2

Flammability: 0 Physical hazard: 0

NFPA ratings

Health: 1

Flammability: 0 Instability: 0

Disclaimer

The information provided in this Material Safety data sheet has been obtained from sources believed to be reliable. Harcros Chemicals Inc provides no warranties, either expressed or implied and assumes no responsibility for the accuracy or completeness of the data contained herein. This information is offered for your information, consideration, and investigation. You should satisfy yourself that you have all current data relevant to your particular use. Harcros Chemicals Inc knows of no medical condition, other than those noted on this material safety data sheet,

which are generally recognized as being aggravated by exposure to this product.

Issue date

05-08-2012

This data sheet contains changes from the previous

version in section(s):

Product and Company Identification: Product and Company Identification Composition / Information on Ingredients: Ingredients

Physical & Chemical Properties: Multiple Properties

Toxicological Information: Toxicological Data

Transport Information: Material Transportation Information

Regulatory Information: United States

Material name: T Det AOS-40 MSDS US 2691 Version #: 01 Revision date: 05-08-2012 Issue date: 05-08-2012 6/6 014nix764y68 fffffeetarper6xf620957:38i25rAMespage 2/010 fax server PRODUCT NAME: HYDROGEN PEROXIDE 35% (ALLGRADES)

MSDS NUMBER:

39247

DATE ISSUED:

09/16/2008

SUPERSEDES:

08/09/2006

ISSUED BY:

008782

Material Safety Data Sheet

1. PRODUCT AND COMPANY IDENTIFICATION

Company

Arkema Inc. 2000 Market Street Philadelphia, Pennsylvania 19103

Oxygenated and Derivatives

Customer Service Telephone Number: (800) 346-7575 (Monday through Friday, 8:30 AM to 5:30 PM EST)

Emergency Information

Transportation:

CHEMTREC: (800) 424-9300 (24 hrs., 7 days a week)

Medical:

Rocky Mountain Poison Center: (303) 623-5716

(24 hrs., 7 days a week)

Product Information

Product name:

HYDROGEN PEROXIDE 35% (ALLGRADES)

Synonyms:

Not available H202

Molecular formula: Chemical family:

peroxides

Molecular weight:

34.01 g/mol

Product use:

Bleaching agent, Oxidizing agent, Cosmetics,

Water treatment

2. HAZARDS IDENTIFICATION

Emergency Overview

Color:

colorless liquid

Physical state: Odor:

pungent

DANGER!

STRONG OXIDIZER.

CONTACT WITH OTHER MATERIAL MAY CAUSE FIRE OR EXPLOSIVE DECOMPOSITION.

CAUSES EYE BURNS.

MAY CAUSE BLINDNESS.

MAY CAUSE RESPIRATORY TRACT IRRITATION.

MAY BE HARMFUL IF SWALLOWED.

Univar Usa FaxServer 7/14/2009 7:38:25 AM PAGE 3/010 Fax Server Potential Health Effects

Primary routes of exposure: Inhalation and skin contact.

Signs and symptoms of acute exposure:

Corrosive to the eyes. May cause irritation of respiratory tract. Effects due to ingestion may include: gastrointestinal symptoms, ulceration, burns, accumulation of fluid in the lungs which may be delayed for several hours.

Skin:

Slightly toxic. Non-irritating. (based on animal studies)

Eyes:

Corrosive. (based on animal studies)

Ingestion:

Practically nontoxic to slightly toxic. (based on animal studies)

3. COMPOSITION/INFORMATION ON INGREDIENTS

Chemical Name	CAS-No.	Wt/Wt	OSHA
			Hazardous
Water	7732-18-5	65 %	N
Hydrogen peroxide (H202)	7722-84-1	35 %	Y

The substance(s) marked with a "Y" in the Hazard column above, are those identified as hazardous chemicals under the criteria of the OSHA Hazard Communication Standard (29 CFR 1910.1200).

This material is classified as hazardous under Federal OSHA regulation.

4. FIRST AID MEASURES

Inhalation:

If inhaled, remove to fresh air. If not breathing, give artificial respiration. If breathing is difficult, give oxygen. Get medical attention.

Skin:

Immediately flush skin with large amounts of water. Remove material from clothing. Wash clothing before reuse. Destroy contaminated shoes.

Eyes:

In case of contact, immediately flush eyes with plenty of water for at least 15 minutes. Get medical attention immediately.

Ingestion:

If swallowed, DO NOT induce vomiting. Get medical attention immediately. If victim is fully conscious, give a cupful of water. Never give anything by mouth to an unconscious person.

Notes to physician:

Exposure to material may cause delayed lung injury resulting in pulmonary edema and pneumonitis. Exposed individuals should be monitored for 72 hours after exposure for the onset of delayed respiratory symptoms.

5. FIRE-FIGHTING MEASURES

Flash point None.

Auto-ignition temperature: not applicable

Lower flammable limit (LFL): not applicable

Upper flammable limit (UFL): not applicable

Univar Usa FaxServer 7/14/2009 7:38:25 AM PAGE 4/010 Fax Server Extinguishing media (suitable): water spray, water fog

Protective equipment:

Fire fighters and others who may be exposed to products of combustion should wear full fire fighting turn out gear (full Bunker Gear) and self-contained breathing apparatus (pressure demand / NIOSH approved or equivalent).

Further firefighting advice:

Fire fighting equipment should be thoroughly decontaminated after use.

Fire and explosion hazards:

Oxidizing Material

Explosive when mixed with combustible material. Avoid breathing fumes from fire exposed material.

6. ACCIDENTAL RELEASE MEASURES

In case of spill or leak:

Stop the leak if you can do so without risk. Ventilate the area. Flush with plenty of water. Avoid contact with cellulose, paper, sawdust or similar substances. Risk of self-ignition or promotion of fires. Combustible materials exposed to hydrogen peroxide should be rinsed immediately with large amounts of water to ensure that all the hydrogen peroxide is removed. Consult a regulatory specialist to determine appropriate state or local reporting requirements, for assistance in waste characterization and/or hazardous waste disposal and other requirements listed in pertinent environmental permits.

7. HANDLING AND STORAGE

Handling

General information on handling:
Do not get in eyes, on skin, or on clothing.
Do not taste or swallow.
Avoid breathing vapor or mist.
Wash thoroughly after handling.
Use only with adequate ventilation.
Avoid contamination.
Keep from contact with clothing and other combustible materials.
Store in tightly closed container.
Emptied container retains vapor and product residue.
Observe all labeled safeguards until container is cleaned, reconditioned or destroyed.
DO NOT CUT, DRILL, GRIND, OR WELD ON OR NEAR THIS CONTAINER.

Storage

General information on storage conditions:

Store away from combustibles and incompatible materials. Store in cool, dry, well ventilated area away from sources of ignition such as flame, sparks and static electricity. Refer to National Fire Protection Association (NFPA) 43A, Code for the Storage of Solid and Liquid Oxidizers.

Storage incompatibility General:

Store separate from acids, alkalies, reducing agents, and combustibles. Store separate from:

Organic materials

Metallic oxides

8. EXPOSURE CONTROLS/PERSONAL PROTECTION

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Hydrogen peroxide (H202) (7722-84-1)

US. ACGIH Threshold Limit Values

Time Weighted Average (TWA): 1 ppm

US. OSHA Table Z-1 Limits for Air Contaminants (29 CFR 1910.1000)

PEL:

1 ppm (1.4 mg/m3)

Only those components with exposure limits are printed in this section. Limits with skin contact designation above have skin contact effect. Air sampling alone is insufficient to accurately quantitate exposure. Measures to prevent significant cutaneous absorption may be required. Limits with a sensitizer designation above mean that exposure to this material may cause allergic reactions.

Engineering controls:

Investigate engineering techniques to reduce exposures below airborne exposure limits. Provide ventilation if necessary to control exposure levels below airborne exposure limits (see above). If practical, use local mechanical exhaust ventilation at sources of air contamination such as open process equipment. Consult ACGIH ventilation manual or NFPA Standard 91 for design of exhaust systems.

Respiratory protection:

Avoid breathing vapor or mist. When airborne exposure limits are exceeded, use NIOSH approved respiratory protection equipment appropriate to the material and/or its components. Full facepiece equipment is recommended and, if used, replaces need for face shield and/or chemical goggles. Consult respirator manufacturer to determine appropriate type equipment for a given application. Observe respirator use limitations specified by NIOSH or the manufacturer. For emergency and other conditions where exposure limits may be significantly exceeded, use an approved full face positive-pressure, self-contained breathing apparatus or positive-pressure airline with auxiliary self-contained air supply. Respiratory protection programs must comply with 29 CFR 1910.134.

Skin protection:

Wear appropriate chemical resistant protective clothing and chemical resistant gloves to prevent skin contact.

When handling this material, gloves of the following type(s) should be worn:

Neoprene

Polyvinylchloride

Impervious butyl rubber gloves

Wear a face shield, chemical goggles and chemical resistant clothing such as an approved splash protective suit made of SBR Rubber, PVC, Gore-Tex or a HAZMAT Splash Protective Suit (Level A, B, or C) when splashing may occur (such as connecting/disconnecting, mechanical first break). For foot protection, wear boots made of NBR, PVC, polyurethane, or neoprene. Overboots made of Latex or PVC, as well as firefighter boots or specialized HAZMAT boots are also permitted. DO NOT wear any form of boot or overboots made of nylon or nylon blends. DO NOT use cotton, wool or leather, as these materials react RAPIDLY with higher concentrations of hydrogen peroxide. Rinse immediately if skin is contaminated. Remove contaminated clothing and shoes immediately. Thoroughly rinse the outside of gloves and protective clothing with water prior to removal. Completely submerge hydrogen peroxide contaminated clothing or other materials in water prior to drying. Residual hydrogen peroxide, if allowed to dry on materials such as paper, fabrics,

coUtiNarlUathErxS@688ro7/lth2009omb38t2Blew cBAGEaus6/010 mafexigerterignite and result in a fire. Clean protective equipment before reuse. Provide a safety shower at any location where skin contact can occur. Wash thoroughly after handling.

Eve protection:

Where there is potential for eye contact, wear a face shield, chemical goggles, and have eye flushing equipment immediately available.

9. PHYSICAL AND CHEMICAL PROPERTIES

Color: Physical state: colorless liquid

Odor:

pungent

pH:

no data available

Density: Vapor pressure: 1.13 g/cm3 (68 deg F (20 deg C)) 24 mmHg (68 deg F (20 deg C))

Relative vapor density: 1.0

Vapor density:

not determined

Boiling point/boiling

226 deg F (108 deg C) range: -27 deg F (-33 deg C)

Freezing point:

completely soluble

Solubility in water: % Volatiles:

100 %

Molecular weight:

34.01 g/mol

Henry's constant:

(Concentration: 50%) 1.00E-02

10. STABILITY AND REACTIVITY

Stability:

This material is chemically stable under normal and anticipated storage, handling and processing conditions.

Materials to avoid:

Metals

Organic materials

Reducing agents Metallic oxides

Bases

Dusts

Combustible materials (e.g., wood, sawdust)

Conditions / hazards to avoid:

Material decomposes with the potential to produce a rupture of unvented closed containers.

Hazardous decomposition products:

This material decomposes if contaminated, causing fire and possible explosions. Oxygen can be liberated at temperatures above ambient.

11. TOXICOLOGICAL INFORMATION

Data on this material and/or its components are summarized below.

Data for HYDROGEN PEROXIDE 35% (ALLGRADES)

Acute toxicity

Practically nontoxic. (rat) LD50 >5000 mg/kg (10%) .

Slightly toxic. (rat) LD50 = 1200 mg/kg (35%).

Dermal:

Slightly toxic. (rat) LD50 >2000 mg/kg (35%).

Skin Irritation:

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                                                    7/010 Fax Server
Non-irritating. (rabbit) (35 %)
 Eye Irritation:
Corrosive. (rabbit) (10 %)
Corrosive. (rabbit) (35 %)
Repeated dose toxicity
Repeated drinking water administration to rat and mouse / affected organ(s):
GI tract / signs: irritation
Repeated inhalation administration to rat and mouse / affected organ(s): nose
/ signs: irritation
Repeated inhalation administration to dog / affected organ(s): upper
respiratory tract, lung / signs:
irritation, emphysema
Chronic oral administration to laboratory animal / affected organ(s): stomach
/ signs: ulceration
Carcinogenicity
Chronic drinking water administration to rat and mouse / affected organ(s):
GI tract / Increased incidence of tumors was reported.
Classified by the International Agency for Research on Cancer as: Group 3:
Unclassifiable as to carcinogenicity in humans.
Genotoxicity
Assessment in Vitro:
Genetic changes were observed in laboratory tests using: bacteria, animal
Cells
Assessment in Vivo:
No genetic changes were observed in laboratory tests using: animals
Human experience
Inhalation:
Throat: irritation. (based on reports of occupational exposure to workers)
Skin contact:
Skin: bleaching of hair. (based on reports of occupational exposure to
workers) Eye contact:
Eye: irritating. (based on reports of occupational exposure to workers)
Ingestion:
GI tract: bloating, ulceration, burns. (accidental exposure to concentrated
solutions) Lung: accumulation of fluid in the lungs, death.
12. ECOLOGICAL INFORMATION
Chemical Fate and Pathway
No data are available.
Ecotoxicology
Data for HYDROGEN PEROXIDE 35% (ALLGRADES)
Aquatic toxicity data:
Slightly toxic. Fish 96 h LC50 between 10 - 37 mg/L
Aquatic invertebrates:
Moderately toxic. Daphnia magna (Water flea) EC50 = 7.7 mg/L
Moderately toxic. Daphnia pulex (Water flea) EC50 = 2.4 mg/L
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Alunivar Usa FaxServer 7/14/2009 7:38:25 AM PAGE 8/010 Fax Server Highly toxic. EC50 = 0.85 mg/L

Microorganisms:

Slightly toxic. Bacteria EC50 = 30 mg/L

13. DISPOSAL CONSIDERATIONS

Waste disposal:

Dilution with water is the preferred method of disposal. Dispose of in accordance with federal, state and local regulations. Consult a regulatory specialist to determine appropriate state or local reporting requirements, for assistance in waste characterization and/or hazardous waste disposal and other requirements listed in pertinent environmental permits. Note: Chemical additions to, processing of, or otherwise altering this material may make this waste management information incomplete, inaccurate, or otherwise inappropriate. Furthermore, state and local waste disposal requirements may be more restrictive or otherwise different from federal laws and regulations.

14. TRANSPORT INFORMATION

US Department of Transportation (DOT)

UN Number

Proper shipping name : Hydrogen peroxide, aqueous solutions

Class : 5.1 Subsidiary hazard class

: (8)

Packaging group : II Marine pollutant

: no

International Maritime Dangerous Goods Code (IMDG)

UN Number : 2014

Proper shipping name : HYDROGEN PEROXIDE, AQUEOUS SOLUTION

Class : 5.1 Subsidiary hazard class : (8) Packaging group : II Marine pollutant : no

15. REGULATORY INFORMATION

Chemical Inventory Status

EU. EINECS EINECS Conforms to

US. Toxic Substances The components of this product are

Control Act all on the TSCA Inventory.

Australia. Industrial AICS

Chemical (Notification and

Assessment) Act

Canada. Canadian All components of this product are on Environmental Protection the Canadian DSL list.

Conforms to

Act (CEPA). Domestic Substances List

(DSL). (Can. Gaz. Part II, Vol. 133)

Japan. Kashin-Hou Law ENCS (JP) Does not conform List

Korea. Toxic Chemical KECI (KR) Conforms to Control Law (TCCL) List

Philippines. The Toxic PICCS (PH) Does not conform Substances and Hazardous

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China. Inventory of

INV (CN)

Does not conform

Existing Chemical Substances

New Zealand. Inventory

NZIOC

Conforms to

of Chemicals (NZIoC), as

published by ERMA New Zealand

United States Federal Regulations

SARA Title III Section 302 Extremely Hazardous Chemicals:

Chemical Name

CAS-No.

SARA

Reportable

Threshold

Hydrogen peroxide (H202) 7722-84-1

Quantities 1000 lbs

Planning Quantity 1000 lbs

SARA Title III - Section 311/312 Hazard Categories: Acute Health Hazard, Fire Hazard, Reactivity Hazard

SARA Title III Section 313 Toxic Chemicals:

SARA 313: This material does not contain any chemical components with known CAS numbers that exceed the threshold (De Minimis) reporting levels established by SARA Title III, Section 313.

Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA) - Reportable Quantity (RQ):

The components in this product are either not CERCLA regulated, regulated but present in negligible concentrations, or regulated with no assigned reportable quantity.

OSHA Regulated Carcinogens (NTP, IARC, OSHA Listed):

No component of this product present at levels greater than or equal to 0.1% is identified as a known or anticipated carcinogen by NTP.

No component of this product present at levels greater than or equal to 0.1% is identified as probable, possible or confirmed human carcinogen by IARC.

No component of this product present at levels greater than or equal to 0.1% is identified as a carcinogen or potential carcinogen by OSHA.

United States State Regulations

Massachusetts Right to Know

Chemical Name

CAS-No.

Hydrogen peroxide (H202)

7722-84-1

Massachusetts Right to Know Extraordinarily Hazardous Substance(s)

Chemical Name

CAS-No.

Hydrogen peroxide (H202)

7722-84-1

New Jersey Right to Know

Chemical Name

CAS-No.

Hydrogen peroxide (H202)

7722-84-1

New Jersey Right to Know

Special Health Hazard Substance(s)

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Hydrogen peroxide (H202)

Pennsylvania Right to Know

Chemical Name Hydrogen peroxide (H202) CAS-No. 7722-84-1

Water

7732-18-5

Pennsylvania Right to Know Environmentally Hazardous Substance(s)

Chemical Name Hydrogen peroxide (H202)

CAS-No. 7722-84-1

California Prop. 65

This product does not contain any chemicals known to the State of California to cause cancer, birth defects, or any other reproductive defects.

16. OTHER INFORMATION

Miscellaneous:

Other information: This MSDS covers the following grades of 35% H202: Albone; Peroxal; Valsterane; AG; B10; CG; CG-HP; CLG; MS; FG; ASG; EG.

----- FOR ADDITIONAL INFORMATION -----CONTACT: MSDS COORDINATOR UNIVAR USA INC. DURING BUSINESS HOURS, PACIFIC TIME (425)889 - 3400----- NOTICE -----

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* * * END OF MSDS * * *

MATERIAL SAFETY DATA SHEET REGULATORY DATA

FILE UPDATE: 07/12/2013

MSDS VERSION DATE: 1-3-2013

PRODUCT CODE: 101082

ORIGINAL ENTRY: 12/26/1990

DESCRIPTION: Sodium Percarbonate

CAS Number and Names of Primary Chemical and/or its Components

15630-89-4 Sodium Percarbonate

100.00

Section 313 Regulated Substances

Regulating Authority

NO Extremely Hazardous Substance subject to section 302 emergency planning and notification requirements (EHS) Hazardous chemical and/or components subject to section 311 and 312 MSDS and inventory requirements (OSH) YES Toxic chemical and/or components subject to toxic chemical release reporting under Section 313 (TOXIC) NO Hazardous contents subject to section 304 spill reporting of Comprehensive Environmental Liability Act (CERCLA) No Subject to the reporting requirements of the EPA Toxicity Characteristic Leaching Process (CFR40 261.24) NO This product, or its components, are listed on or are exempt from the Toxic Substance Control Act (TSCA) YES Contains a Toxic Air Pollutant listed under the 1990 Clean Air Act Ammendments [42 USC sec 7412(b)(1)] NO Subject to the EPA Risk Management Program under Section 112(r) of the Clean Air Act and 40 CFR Part 68 NO

SARA Title III Hazard Categories

Department of Transportation Data

Fire Hazard	NO	Sodium carbonate per	oxyhydrate
Reactivity Hazard	NO	Hazard Class:	5.1
Sudden Release of Pressure	NO	ID Number:	UN3378
Acute (immediate health hazard)	YES	ERG Guide #:	140
Chronic (delayed health hazard)	NO	Packing Group:	II
Extremely Hazardous Substance	NO	RQ Pounds:	

HMIS & NFPA Hazard Ratings

H Health 1
M Flammability 1
I Reactivity 2
S PPE

N Health

F Flammability
P Reactivity

Special

2008 EMERGENCY RESPONSE GUIDE SHEET

POTENTIAL HAZARDS

Guide Number: 140

FIRE OR EXPLOSION

- These substances will accelerate burning when involved in a fire.
- Some may decompose explosively when heated or involved in a fire.
- May explode from heat or contamination.
- Some will react explosively with hydrocarbons (fuels).
- May ignite combustibles (wood, paper, oil, clothing, etc.).
- · Containers may explode when heated.
- Runoff may create fire or explosion hazard.

HEALTH

- Inhalation, ingestion or contact (skin, eyes) with vapors or substance may cause severe injury, burns or death.
- Fire may produce irritating, corrosive and/or toxic gases.
- Runoff from fire control or dilution water may cause pollution.

PUBLIC SAFETY

- CALL Emergency Response Telephone Number on Shipping Paper first. If Shipping Paper not available or no answer, refer to appropriate telephone number listed on the inside back cover.
- As an immediate precautionary measure, isolate spill or leak area in all directions for at least 50 meters (150 feet) for liquids and at least 25 meters (75 feet) for solids.
- · Keep unauthorized personnel away.
- Stay upwind.
- · Keep out of low areas.
- · Ventilate closed spaces before entering.

PROTECTIVE CLOTHING

- Wear positive pressure self-contained breathing apparatus (SCBA).
- Wear chemical protective clothing that is specifically recommended by the manufacturer. It may provide little or no thermal protection.
- Structural firefighters' protective clothing will only provide limited protection.

EVACUATION

Large Spill

Consider initial downwind evacuation for at least 100 meters (330 feet).

Fire

 \bullet If tank, rail car or tank truck is involved in a fire, ISOLATE for 800 meters (1/2 mile) in all directions; also, consider initial evacuation for 800 meters (1/2 mile) in all directions.

EMERGENCY RESPONSE

FIRE

Small Fire

• Use water. Do not use dry chemicals or foams. CO2 or Halon® may provide limited control.

Large Fire

- Flood fire area with water from a distance.
- Do not move cargo or vehicle if cargo has been exposed to heat.
- Move containers from fire area if you can do it without risk.

2008 EMERGENCY RESPONSE GUIDE SHEET

Fire involving Tanks or Car/Trailer Loads

- Fight fire from maximum distance or use unmanned hose holders or monitor nozzles.
- Cool containers with flooding quantities of water until well after fire is out.
- ALWAYS stay away from tanks engulfed in fire.
- For massive fire, use unmanned hose holders or monitor nozzles; if this is impossible, withdraw from area and let fire burn.

Guide Number: 140

SPILL OR LEAK

- Keep combustibles (wood, paper, oil, etc.) away from spilled material.
- Do not touch damaged containers or spilled material unless wearing appropriate protective clothing.
- Stop leak if you can do it without risk.
- Do not get water inside containers.

Small Dry Spill

• With clean shovel place material into clean, dry container and cover loosely; move containers from spill area.

Small Liquid Spill

• Use a non-combustible material like vermiculite or sand to soak up the product and place into a container for later disposal.

Large Spill

- Dike far ahead of liquid spill for later disposal.
- Following product recovery, flush area with water.

FIRST AID

- Move victim to fresh air.
- Call 911 or emergency medical service.
- Give artificial respiration if victim is not breathing.
- Administer oxygen if breathing is difficult.
- Remove and isolate contaminated clothing and shoes.
- Contaminated clothing may be a fire risk when dry.
- In case of contact with substance, immediately flush skin or eyes with running water for at least 20 minutes.
- Keep victim warm and quiet.
- Ensure that medical personnel are aware of the material(s) involved and take precautions to protect themselves.



20 Buckingham Court, Cartersville, GA 30120 770-386-3555/Fax 770-386-2009

MATERIAL SAFETY DATA SHEET SODIUM CARBONATE PEROXYHYDRATE



MSDS No. 001

SECTION 1: PRODUCT AND COMPANY IDENTIFICATION

SECTION 2: COMPOSITION/INFORMATION ON INGREDIENTS

SECTION 3: HAZARDS IDENTIFICATION

SECTION 4: FIRST AID MEASURES

SECTION 5: FIRE FIGHTING MEASURES

SECTION 6: ACCIDENTAL RELEASE MEASURES

SECTION 7: HANDLING AND STORAGE

SECTION 8: EXPOSURE CONTROLS/PERSONAL PROTECTION

SECTION 9: PHYSICAL AND CHEMICAL PROPERTIES

SECTION 10: STABILITY AND REACTIVITY

SECTION 11: TOXICOLOGICAL INFORMATION

SECTION 12: ECOLOGICAL INFORMATION

SECTION 13: DISPOSAL CONSIDERATION

SECTION 14: TRANSPORT INFORMATION

SECTION 15: REGULATORY INFORMATION

SECTION 16: ADDITIONAL INFORMATION SECTION 17: LABEL INFORMATION

SECTION 1: PRODUCT AND COMPANY IDENTIFICATION



SECTION 2: COMPOSITION/INFORMATION ON INGREDIENTS

INGREDIENT	%CONC.	C.A.S.	ACGIH-TLV	OSHA-PEL
SODIUM CARBONATE PEROXYHYDRATE	98-100	15630-89-4	10 MG/M3 TWA INHALABLE PARTICLES	15 MG/M3 TWA TOTAL DUST
			3 MG/M3 TWA RESPIRABLE PARTICULATE	5 MG/M3 TWA RESPIRABLE FRACTION
			PARTICLES NOT OTHERWISE SPECIFIED (PNOC)	PARTICLES NOT OTHERWISE REGULATED (PNOR)

SECTION 3: HAZARDS IDENTIFICATION

EMERGENCY OVERVIEW.

PRODUCT IS WHITE HYGROSCOPIC CRYSTALS.

OXIDIZER. MAY BE HARMFUL OR FATAL IF SWALLOWED

OR INHALED. MAY CAUSE SEVERE EYE AND

RESPIRATORY TRACT IRRITATION OR BURNS. MAY

CAUSE SKIN IRRITATION. CONTACT WITH OTHER

MATERIALS MAY CAUSE FIRE.

POTENTIAL HEALTH EFFECTS:

INGESTION.

VOMITING AND DIARRHEA.

SKIN CONTACT.

MAY CAUSE SKIN IRRITATION AND REDNESS.

INHALATION.

IRRITATING TO THE RESPIRATORY TRACT.

COUGHING, SNEEZING, DIFFICULTY BREATHING AND

SORE THROAT.

EYE CONTACT.

MAY CAUSE IRRITATION TO THE EYES, INCLUDING

PAIN. REDNESS AND REVERSIBLE DAMAGE.

OUDION OUTDOWN IF LEUCY HITDUATE

AGGRAVATED	PERSONS	WITH	CHRONIC	RESPIRATORY	OR	SKIN
	DISEASES	3.				
SUBCHRONIC (TARGET ORGAN	1)					
EFFECTS	ATAG OM					
CHRONIC EFFECTS	NO DATA			1		
CARCINOGENICITY						
NTP	NO					
IARC	NO					
OSHA	NO					
OTHER TOXICOLOGICAL DATA	NO DATA					
PRINCIPAL ROUTES OF ENTE	RYEYE CONT	CACT				
	SKIN CON	TACT				
	INHALATI	ON				

SECTION 4: FIRST AID MEASURES

EYE CONTACTWASH EYES IMMEDIATELY WITH LARGE AMOUNTS OF
WATER OR NORMAL SALINE, OCCASIONALLY LIFTING
UPPER AND LOWER LIDS, UNTIL NO EVIDENCE OF
CHEMICAL REMAINS (AT LEAST 15-20 MINUTES). GET
MEDICAL ATTENTION IMMEDIATELY.
SKIN CONTACTREMOVE CONTAMINATED CLOTHING AND SHOES
IMMEDIATELY. WASH WITH SOAP OR MILD DETERGENT
AND LARGE AMOUNTS OF WATER UNTIL NO EVIDENCE OF
CHEMICAL REMAINS (AT LEAST 15-20 MINUTES). GET
MEDICAL ATTENTION IMMEDIATELY.
INHALATIONIMMEDIATELY REMOVE VICTIM TO FRESH AIR, IF
VICTIM IS NOT BREATHING A QUALIFIED PERSON
SHOULD IMMEDIATELY BECIN ARTIFICIAL
RESPIRATION, OBTAIN IMMEDIATE MEDICAL
ASSISTANCE !
INGESTIONOBTAIN IMMEDIATE MEDICAL ASSISTANCE - DO NOT
INDUCE VOMITING. IF VOMITING OCCURS
SPONTANEOUSLY, KEEP HEAD BELOW HIPS TO AVOID
ASPIRATION.
NOTE TO PHYSICIANTREAT SYMPTOMATICALLY

SECTION 5: FIRE FIGHTING MEASURES

FLASH POINTNON	E		
METHODNOT	APPLICABLE		
IGNITION TEMPERATURENOT	APPLICABLE	1	
UPPER FLAMMABLE LIMITNOT	APPLICABLE		
LOWER FLAMMABILITY LIMITNOT	APPLICABLE		
SENSITIVITY TO MECHANICAL			
IMPACTNO			
SENSITIVITY TO STATIC			
DISCHARGENOT	EXPECTED		
EXTINGUISHING MEDIAUSE	SUITABLE EXTINGUISHING	AGENTS	FOR

3

SURROUNDING FIRE.

SPECIAL FIREFIGHTING

PROCEDURES...... EVACUATE AREA AND FIGHT FIRE FROM A SAFE

DISTANCE. AS IN ANY FIRE, WEAR SELF-CONTAINED BREATHING APPARATUS PRESSURE-DEMAND, AND FULL

PROTECTIVE GEAR.

FLAMMABILITY CLASS (OSHA) . . . NOT APPLICABLE

SECTION 6: ACCIDENTAL RELEASE MEASURES

ACTION TO BE TAKEN FOR

SPILLS OR RELEASES......DO NOT TOUCH SPILLED MATERIAL. STOP FLOW OF

MATERIAL, IF THIS IS WITHOUT RISK. PREVENT MATERIAL FROM ENTERING SEWERS AND WATERWAYS.

ENSURE CLEAN-UP IS CONDUCTED BY TRAINED PERSONNEL. WEAR APPROPRIATE PERSONAL

PROTECTIVE EQUIPMENT.

SHOVEL INTO CLEAN, DRY LABELED CONTAINERS. USE

EXTREME CARE TO PREVENT CONTAMINATION WITH

COMBUSTIBLE OR ORGANIC MATERIALS.

KEEP UNNECESSARY PERSONNEL AWAY. CLOSE OFF

AREA.

SECTION 7: HANDLING AND STORAGE

HANDLING PROCEDURES......AVOID GETTING IN EYES, ON SKIN OR ON CLOTHING.

AVOID BREATHING FUMES OR DUST FROM THIS

MATERIAL. AVOID GENERATION OF MIST OR DUST. USE THIS PRODUCT ONLY WITH ADEQUATE VENTILATION. PREVENT CONTACT WITH COMBUSTIBLE OR ORGANIC

MATERIALS. LABEL CONTAINERS AND KEEP THEM TIGHTLY CLOSED WHEN NOT IN USE. WASH THOROUGHLY

AFTER HANDLING.

STORAGE PROCEDURES......STORE IN TIGHTLY CLOSED CONTAINERS IN A CLEAN,

DRY, WELL-VENTILATED AREA. STORE AWAY FROM SOURCES OF HEAT, INCOMPATIBLE MATERIALS, AND

COMBUSTIBLE MATERIALS.

SECTION 8: EXPOSURE CONTROLS/PERSONAL PROTECTION

ENGINEERING CONTROLS......DESIGN AND IMPLEMENT CONTROLS AND PROCEDURES TO AVOID ACCUMULATION OF DUST.

6

RESPIRATORY PROTECTION	SEE SECTION 2 - SPECIFIC INGREDIENTS THE SPECIFIC RESPIRATOR SELECTED MUST BE BASED ON CONTAMINATION LEVELS FOUND IN THE WORK PLACE, MUST NOT EXCEED THE WORKING LIMITS OF THE RESPIRATOR AND BE JOINTLY APPROVED BY THE NATIONAL INSTITUTE FOR OCCUPATIONAL SAFETY AND HEALTH AND THE MINE SAFETY AND HEALTH ADMINISTRATION (NIOSH-MSHA). IMPERVIOUS GLOVES AND CLOTHING TO PREVENT SKIN CONTACT.
	DUST OR SPLASH PROOF CHEMICAL SAFETY GOGGLES OR
	FACE SHIELDS (EIGHT-INCH MINIMUM). WHERE THERE IS ANY POSSIBILITY THAT AN EMPLOYEE'S EYES MAY BE EXPOSED TO THIS SUBSTANCE, THE EMPLOYER SHOULD PROVIDE AN EYE WASH FOUNTAIN WITHIN THE IMMEDIATE WORK AREA FOR EMERGENCY USE. WHERE THERE IS ANY POSSIBILITY THAT AN EMPLOYEE'S SKIN MAY BE EXPOSED TO THIS SUBSTANCE, THE EMPLOYER SHOULD PROVIDE A QUICK DRENCH SHOWER WITHIN THE IMMEDIATE WORK AREA FOR EMERGENCY USE. EMPLOYEES SHOULD BE TRAINED IN EMERGENCY MEASURES IN CASE OF LEAK, SPILL, FIRE OR EXPLOSION. TRAINING IN USE OF EMERGENCY RESPONSE EQUIPMENT SHOULD BE PROVIDED TO INCLUDE LOCATION OF PHONE, FIRST AID KITS,
	ALARMS, FIRE EXTINGUISHERS FOR THE SPECIFIC WORKPLACE. USE LOCAL EXHAUST AND PROCESS ENCLOSURE TO CONTROL DUSTS OR VAPORS. USE OF A CORROSION-RESISTANT VENTILATION SYSTEM IS RECOMMENDED.

SECTION 9: PHYSICAL AND CHEMICAL PROPERTIES

VAPOR DENSITY (AIR=1).....NO DATA FREEZING/MELTING POINT.....NO DATA PHYSICAL STATE.....SOLID ODOR.....ODORLESS COLOR......WHITE CRYSTALLINE ODOR THRESHOLD (PPM).....NO DATA VOLATILES % VOLUME......NOT APPLICABLE EVAPORATION RATE (BUTYL ACETATE=1.0).....NOT APPLICABLE SPECIFIC GRAVITY (WATER=1.0).....0.97 MOLECULAR WEIGHT......324.0 ACID/ALKALINITY (MEQ/G)..., UNKNOWN VOC (EPA METHOD 24).....NOT APPLICABLE SOLUBILITY IN ORGANIC SOLVENTS......NO DATA BOILING POINT........NO DATA VAPOR PRESSURE MM/HG

MATERIAL PARTIES AND ADMINISTRATION ADMINISTRATION AND ADMINISTRATION			
5			

OUDION OUTDOMNIT I FUNCTION IN THE

(20°C)							•	8		NO.	DATE
SOLUBILITY											
OCTANOL/WAT	CER	P	AR	TI	T	IC	N				
COEFFICIEN					•			4		NO.	DATE

SECTION 10: STABILITY AND REACTIVITY

STABILITY......STABLE AS SOLID

HAZARDOUS POLYMERIZATION...WILL NOT OCCUR

HAZARDOUS THERMAL

DECOMPOSITION/COMBUSION

PRODUCTS......OXYGEN

INCOMPATIBILITY

(MATERIALS TO AVOID)....BASES

FLAMMABLE AND COMBUSTIBLE MATERIALS, NITROGEN

COMPOUNDS, ACIDS, METAL OXIDES AND MOISTURE.

CONDITIONS TO AVOID....CONTACT WITH INCOMPATIBLE SUBSTANCES.

EXPOSURE TO INTENSE HEAT

MOISTURE

HIGH TEMPERATURES

AVOID CONTACT WITH ALL COMBUSTIBLE MATERIALS.

SECTION 11: TOXICOLOGICAL INFORMATION

LD50	ORAL240	00 mg/kg	RAT				
	220	00 mg/kg	MOUSE				
LD50	DERMALNO	DATA					
LC50	INHALATIONNO	DATA					
	R						
AMES	TEST RESULTSNO	DATA AV	AILABLE	COVERING	AMES	TEST	RESULTS

SECTION 12: ECOLOGICAL INFORMATION

ECOTOXICOLOGICAL
INFORMATION.........................NO DATA AT THIS TIME
CHEMICAL FATE INFORMATION...NO DATA AT THIS TIME

SECTION 13: DISPOSAL CONSIDERATION

6

DISPOSAL METHODIN	ACCORDANCE WITH FEDERAL, STATE, LOCAL
R	GULATIONS
REPORTABLE QUANTITYMA	Y MEET THE REQUIREMENTS OF IGNITABLE WASTE
)C	01.

SECTION 14: TRANSPORT INFORMATION

DOT SIJIPPING NAME	******	
	SODIUM CARBONATE PE	ROXYHYDRATE
DOT HAZARD CLASS	5.1	
DOT LABELS	OXIDIZER	
UN NUMBER	UN 3378	
PLACARDS		
IATA	NOT REGULATED	
IMO/IMDG	NOT REGULATED	
TRANSPORT EMERGENCY		
PROCEDURES	CONTACT CHEMTREC 1-	300-424-9300
OTHER	NOT APPLICABLE	

SECTION 15: REGULATORY INFORMATION

SARA SECTION 302YES
SARA (313) CHEMICALSNO
EPA TSCA INVENTORYAPPEARS
CERCLA SECTION 103NO
CANADIAN WHMIS
CLASSIFICATION
CANADIAN DOMESTIC
SUBSTANCES LIST (DSL)APPEARS
CALIFORNIA PROPOSITION 65NO
EINECS INVENTORYAPPEARS OTHERTHIS PRODUCT IS REGULATED BY THE FOLLOWING STATES:
CA; FL; MA; MN; NJ; PA

SECTION 16: ADDITIONAL INFORMATION

FROM	MATERIAL SAFETY DATA SHEET WAS PRODUCED RELIABLE SOURCES. HOWEVER, I'T IS PROVIDED
WITH	OUT REPRESENTATION OR WARRANTY EXPRESSED OR IED REGARDING ACCURACY OR CORRECTNESS.
COND	ITIONS AND METHODS OF USE ARE BEYOND THE ROL AND KNOWLEDGE OF CHEMICAL SPECIALTY GROUP, INC.

7

CHEMICAL SPECIALTY GROUP, INC. DOES NOT ASSUME ANY

RESPONSIBILITY AND EXPRESSLY DISCLAIMS

LIABILITY FOR INJURY, LOSS, DAMAGE OR EXPENSES

ARISING FROM THE USE OF THIS PRODUCT.

COLIGINAL COMPANIES OF CONTINUES OF

ABBREVIATIONS USED......ABBREVIATIONS USED THROUGHOUT THIS MSDS ARE:

ACGIH = AMERICAN CONFERENCE OF GOVERNMENTAL

INDUSTRIAL HYGIENISTS

TWA = TIME WEIGHTED AVERAGE (EXPOSURE VALUES)

STEL = SHORT TERM EXPOSURE LIMITS

OSHA = OCCUPATIONAL SAFETY AND HEALTH

ADMINISTRATION

PEL = PERMITTED EXPOSURE LIMITS

ppm = PARTS PER MILLION

mg = MILLIGRAMS

NIOSH = NATIONAL INSTITUTE FOR OCCUPATIONAL

HEALTH AND SAFETY

MSHA = MINE SAFETY AND HEALTH ADMINISTRATION

lb = POUNDS

m3 = PER METRE CUBED

NTP = NATIONAL TOXICOLOGICAL PROGRAM

g = GRAMS

ml = MILLILITRE

RTECS = REGISTRY OF TOXICS EFFECTS OF CHEMICAL

SUBSTANCES (NIOSH)

SECTION 17: LABEL INFORMATION

WARNING/PRECAUTIONS......PRODUCT IS WHITE HYGROSCOPIC CRYSTALS.

OXIDIZER, MAY BE HARMFUL OR FATAL IF SWALLOWED

OR INHALED. MAY CAUSE SEVERE EYE AND

RESPIRATORY TRACT IRRITATION OR BURNS. MAY

CAUSE SKIN IRRITATION. CONTACT WITH OTHER

MATERIALS MAY CAUSE FIRE.

FOR FURTHER INFORMATION....SEE THE MATERIAL SAFETY DATA SHEET

SUPPLIER..... CHEMICAL SPECIALTY GROUP, INC.

20 BUCKINGHAM COURT

CARTERSVILLE, GA 30120 TELE 770-386-3555 FAX 770-386-2009

MATERIAL SAFETY DATA SHEET

KlozürTM



MSDS Ref. No.: 7775-27-1-12 Date Approved: 02/22/2005

Revision No.: 1

This document has been prepared to meet the requirements of the U.S. OSHA Hazard Communication Standard, 29 CFR 1910.1200; the Canada's Workplace Hazardous Materials Information System (WHMIS) and, the EC Directive, 2001/58/EC.

1. PRODUCT AND COMPANY IDENTIFICATION

PRODUCT NAME:

Klozür™

SYNONYMS:

Sodium Persulfate, Sodium Peroxydisulfate; Disodium

Peroxydisulfate

GENERAL USE:

In situ and ex situ chemical oxidation of contaminants and

compounds of concern for environmental remediation applications.

MANUFACTURER

EMERGENCY TELEPHONE NUMBERS

FMC CORPORATION Active Oxidants Division 1735 Market Street Philadelphia, PA 19103 (215) 299-6000 (General Information) (800) 424-9300 (CHEMTREC - U.S.) (303) 595-9048 (Medical - Call Collect)

2. HAZARDS IDENTIFICATION

EMERGENCY OVERVIEW:

- White, odorless, crystals
- Oxidizer,
- Decomposes in storage under conditions of moisture (water/water vapor) and/or excessive heat causing release of oxides of sulfur and oxygen that supports combustion. Decomposition could form a high temperature melt. See Section 10 ("Stability and Reactivity").

POTENTIAL HEALTH EFFECTS: Airborne persulfate dust may be irritating to eyes, nose, lungs, throat and skin upon contact. Exposure to high levels of persulfate dust may cause difficulty in breathing in sensitive persons.

Page 1 of 9

3. COMPOSITION / INFORMATION ON INGREDIENTS

Chemical Name	CAS#	Wt.%	EC No.	EC Class
Sodium Persulfate	7775-27-1	>99	231-892-1	Not classified as hazardous

4. FIRST AID MEASURES

EYES: Flush with plenty of water. Get medical attention if irritation occurs and persists.

SKIN: Wash with plenty of soap and water. Get medical attention if irritation occurs and persists.

INGESTION: Rinse mouth with water. Dilute by giving 1 or 2 glasses of water. Do not induce vomiting. Never give anything by mouth to an unconscious person. See a medical doctor immediately.

INHALATION: Remove to fresh air. If breathing difficulty or discomfort occurs and persists, contact a medical doctor.

NOTES TO MEDICAL DOCTOR: This product has low oral toxicity and is not irritating to the eyes and skin. Flooding of exposed areas with water is suggested, but gastric lavage or emesis induction for ingestions must consider possible aggravation of esophageal injury and the expected absence of system effects. Treatment is controlled removal of exposure followed by symptomatic and supportive care.

5. FIRE FIGHTING MEASURES

EXTINGUISHING MEDIA: Deluge with water.

FIRE / EXPLOSION HAZARDS: Product is non-combustible. On decomposition releases oxygen which may intensify fire. Presence of water accelerates decomposition.

FIRE FIGHTING PROCEDURES: Do not use carbon dioxide or other gas filled fire extinguishers; they will have no effect on decomposing persulfates. Wear full protective clothing and self-contained breathing apparatus.

FLAMMABLE LIMITS: Non-combustible

SENSITIVITY TO IMPACT: No data available

Page 2 of 9

SENSITIVITY TO STATIC DISCHARGE: Not available

6. ACCIDENTAL RELEASE MEASURES

RELEASE NOTES: Spilled material should be collected and put in approved DOT container and isolated for disposal. Isolated material should be monitored for signs of decomposition (fuming/smoking). If spilled material is wet, dissolve with large quantity of water and dispose as a hazardous waste. All disposals should be carried out according to regulatory agencies procedures.

7. HANDLING AND STORAGE

HANDLING: Use adequate ventilation when transferring product from bags or drums. Wear respiratory protection if ventilation is inadequate or not available. Use eye and skin protection. Use clean plastic or stainless steel scoops only.

STORAGE: Store (unopened) in a cool, clean, dry place away from point sources of heat, e.g. radiant heaters or steam pipes. Use first in, first out storage system. Avoid contamination of opened product. In case of fire or decomposition (fuming/smoking) deluge with plenty of water to control decomposition. For storage, refer to NFPA Bulletin 430 on storage of liquid and solid oxidizing materials.

COMMENTS: VENTILATION: Provide mechanical general and/or local exhaust ventilation to prevent release of dust into work environment. Spills should be collected into suitable containers to prevent dispersion into the air.

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

EXPOSURE LIMITS

Chemical Name	ACGIH	OSHA	Supplier
Sodium Persulfate	0.1 mg/m ³ (TWA)		

ENGINEERING CONTROLS: Provide mechanical local general room ventilation to prevent release of dust into the work environment. Remove contaminated clothing immediately and wash before reuse.

PERSONAL PROTECTIVE EQUIPMENT

EYES AND FACE: Use cup type chemical goggles. Full face shield may be used.

RESPIRATORY: Use approved dust respirator when airborne dust is expected.

Page 3 of 9

PROTECTIVE CLOTHING: Normal work clothes. Rubber or neoprene footwear.

GLOVES: Rubber or neoprene gloves. Thoroughly wash the outside of gloves with soap and water prior to removal. Inspect regularly for leaks.

9. PHYSICAL AND CHEMICAL PROPERTIES

ODOR:

None

APPEARANCE:

White crystals

AUTOIGNITION TEMPERATURE:

Not applicable. No evidence of combustion up to 800°C.

Decomposition will occur upon heating.

BOILING POINT:

Not applicable

COEFFICIENT OF OIL / WATER:

Not applicable

DENSITY / WEIGHT PER VOLUME:

Not available

EVAPORATION RATE:

Not applicable (Butyl Acetate = 1)

FLASH POINT:

Non-combustible

MELTING POINT:

Decomposes

ODOR THRESHOLD:

Not applicable

OXIDIZING PROPERTIES:

Oxidizer

PERCENT VOLATILE:

Not applicable

pH:

typically 5.0 - 7.0 @ 25 °C (1% solution)

SOLUBILITY IN WATER:

73 % @ 25 °C (by wt.)

SPECIFIC GRAVITY:

 $2.6 (H_2O=1)$

VAPOR DENSITY:

Not applicable (Air = 1)

VAPOR PRESSURE:

Not applicable

10. STABILITY AND REACTIVITY

CONDITIONS TO AVOID:

Heat, moisture and contamination.

STABILITY:

Stable (becomes unstable in presence of heat,

moisture and/or contamination).

POLYMERIZATION:

Will not occur

INCOMPATIBLE MATERIALS:

Acids, alkalis, halides (fluorides, chlorides,

bromides and iodides), combustible materials, most metals and heavy metals, oxidizable materials, other oxidizers, reducing agents, cleaners, and organic or carbon containing compounds. Contact

Page 4 of 9

> with incompatible materials can result in a material decomposition or other uncontrolled reactions.

HAZARDOUS DECOMPOSITION PRODUCTS: Oxygen that supports combustion and oxides of sulfur.

COMMENTS: PRECAUTIONARY STATEMENT: Pumping and transport of Klozür persulfate requires appropriate precautions and design considerations for pressure and thermal relief.

Decomposing persulfates will evolve large volumes of gas and/or vapor, can accelerate exponentially with heat generation, and create significant and hazardous pressures if contained and not properly controlled or mitigated.

Use with alcohols in the presence of water has been demonstrated to generate conditions that require rigorous adherence to process safety methods and standards to prevent escalation to an uncontrolled reaction.

11. TOXICOLOGICAL INFORMATION

EYE EFFECTS: Non-irritating (rabbit) [FMC Study Number: ICG/T-79.029]

SKIN EFFECTS: Non-irritating (rabbit) [FMC Study Number: ICG/T-79.029]

DERMAL LD₅₀: > 10 g/kg [FMC Study Number: ICG/T-79.029]

ORAL LD₅₀: 895 mg/kg (rat) [FMC Study Number: ICG/T-79.029]

INHALATION LC₅₀: 5.1 mg/l (rat) [FMC 195-2017]

SENSITIZATION: May be sensitizing to allergic persons. [FMC Study Number: ICG/T-79.029]

TARGET ORGANS: Eyes, skin, respiratory passages

ACUTE EFFECTS FROM OVEREXPOSURE: Dust may be harmful and irritating. May be harmful if swallowed.

CHRONIC EFFECTS FROM OVEREXPOSURE: Sensitive persons may develop dermatitis and asthma [Respiration 38:144, 1979]. Groups of male and female rats were fed 0, 300 or 3000 ppm sodium persulfate in the diet for 13 weeks, followed by 5000 ppm for 5 weeks. Microscopic examination of tissues revealed some injury to the gastrointestinal tract at the high dose (3000 ppm) only. This effect is not unexpected for an oxidizer at high concentrations. [Ref. FMC I90-1151, Toxicologist 1:149, 1981].

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CARCINOGENICITY:

NTP:

Not listed

IARC:

Not listed

OSHA:

Not listed

OTHER:

ACGIH: Not listed

12. ECOLOGICAL INFORMATION

ECOTOXICOLOGICAL INFORMATION:

Bluegill sunfish, 96-hour $LC_{50} = 771 \text{ mg/L}$ [FMC Study I92-1250] Rainbow trout, 96-hour $LC_{50} = 163 \text{ mg/L}$ [FMC Study I92-1251] Daphnia, 48-hour $LC_{50} = 133 \text{ mg/L}$ [FMC Study I92-1252] Grass shrimp, 96-hour $LC_{50} = 519$ mg/L [FMC Study I92-1253]

CHEMICAL FATE INFORMATION: Biodegradability does not apply to inorganic substances.

13. DISPOSAL CONSIDERATIONS

DISPOSAL METHOD: Dispose as a hazardous waste in accordance with local, state and federal regulatory agencies.

14. TRANSPORT INFORMATION

U.S. DEPARTMENT OF TRANSPORTATION (DOT)

PROPER SHIPPING NAME:

Sodium Persulfate

PRIMARY HAZARD CLASS / DIVISION:

5.1 (Oxidizer)

UN/NA NUMBER:

UN 1505

PACKING GROUP:

III

LABEL(S):

5.1 (Oxidizer)

PLACARD(S):

5.1 (Oxidizer)

MARKING(S):

Sodium Persulfate, UN 1505

ADDITIONAL INFORMATION:

Hazardous Substance/RQ: Not applicable

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49 STCC Number: 4918733

This material is shipped in 225 lb. fiber drums, 55 lb. poly bags and 1000 - 2200 lb. IBC's (supersacks).

INTERNATIONAL MARITIME DANGEROUS GOODS (IMDG)

PROPER SHIPPING NAME:

Sodium Persulfate

INTERNATIONAL CIVIL AVIATION ORGANIZATION (ICAO) / INTERNATIONAL AIR TRANSPORT ASSOCIATION (IATA)

PROPER SHIPPING NAME:

Sodium Persulfate

OTHER INFORMATION:

Protect from physical damage. Do not store near acids, moisture or heat.

15. REGULATORY INFORMATION

UNITED STATES

SARA TITLE III (SUPERFUND AMENDMENTS AND REAUTHORIZATION ACT)

SECTION 302 EXTREMELY HAZARDOUS SUBSTANCES (40 CFR 355, APPENDIX A): Not applicable

SECTION 311 HAZARD CATEGORIES (40 CFR 370):

Fire Hazard, Immediate (Acute) Health Hazard

SECTION 312 THRESHOLD PLANNING QUANTITY (40 CFR 370):

The Threshold Planning Quantity (TPQ) for this product, if treated as a mixture, is 10,000 lbs; however, this product contains the following ingredients with a TPQ of less than 10,000 lbs.: None

SECTION 313 REPORTABLE INGREDIENTS (40 CFR 372):

Not listed

CERCLA (COMPREHENSIVE ENVIRONMENTAL RESPONSE COMPENSATION AND LIABILITY ACT)

CERCLA DESIGNATION & REPORTABLE QUANTITIES (RQ) (40 CFR 302.4): Unlisted, RQ = 100 lbs., Ignitability

TSCA (TOXIC SUBSTANCE CONTROL ACT)
TSCA INVENTORY STATUS (40 CFR 710):

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20			

Date: 02/22/2005 KlozürTM (7775-27-1-12)

Listed

RESOURCE CONSERVATION AND RECOVERY ACT (RCRA) RCRA IDENTIFICATION OF HAZARDOUS WASTE (40 CFR 261):

Waste Number: D001

CANADA

WHMIS (WORKPLACE HAZARDOUS MATERIALS INFORMATION SYSTEM):

Product Identification Number: 1505

Hazard Classification / Division: Class C (Oxidizer), Class D, Div. 2, Subdiv. B. (Toxic)

Ingredient Disclosure List:

Listed

INTERNATIONAL LISTINGS

Sodium persulfate:

Australia (AICS): Listed

China: Listed

Japan (ENCS): (1)-1131

Korea: KE-12369

Philippines (PICCS): Listed

HAZARD, RISK AND SAFETY PHRASE DESCRIPTIONS:

EC Symbols:

(Not classified as hazardous)

EC Risk Phrases:

(Not classified as hazardous)

EC Safety Phrases:

(Not classified as hazardous)

16. OTHER INFORMATION

HMIS

Iealth	1
lammability	0
hysical Hazard	1
ersonal Protection (PPE)	J
	J

Protection = J (Safety goggles, gloves, apron & combination dust & vapor respirator)

HMIS = Hazardous Materials Identification System

Degree of Hazard Code:

4 = Severe

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KlozürTM (7775-27-1-12)

- 3 = Serious
- 2 = Moderate
- 1 = Slight
- 0 = Minimal

NFPA

Health	1
Flammability	0
Reactivity	1
Special	OX

SPECIAL = OX (Oxidizer)

NFPA = National Fire Protection Association

Degree of Hazard Code:

- 4 = Extreme
- 3 = High
- 2 = Moderate
- 1 = Slight
- 0 = Insignificant

REVISION SUMMARY:

New MSDS

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Date: 02/22/2005

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SAFETY DATA SHEET

North American Version

IXPER(R) 70C CALCIUM PEROXIDE GRANULES

1.1. Identification of the substance or preparation

Product name

IXPER(R) 70C CALCIUM PEROXIDE GRANULES

Molecular Weight

: 72.1 g/mol

1.2. Use of the Substance/Preparation

Recommended use

Oxidising Agents

Chemical industry - Agriculture industry

Soil remediation

For further information, please contact: Supplier

1.3. Company/Undertaking Identification

Address

: SOLVAY CHEMICALS, INC.

3333 RICHMOND AVENUE HOUSTON TX 77098-3099

United States

1.4. Emergency and contact telephone numbers

Emergency telephone

: 1 (800) 424-9300 CHEMTREC ® (USA & Canada)

01-800-00-214-00 (MEX. REPUBLIC)

(product information):

Contact telephone number : US: +1-800-765-8292 (Product information) US: +1-713-525-6500 (Product information)

2.1. Emergency Overview:

NFPA

: H= 2 F= 0 I= 1 S= OX

HMIS

: H= 2 F= 0 R= 1 PPE = Supplied by User; dependent on local

conditions

General Information

Appearance

: Granular

Colour

: light yellow : odourless

Odour

Main effects - The preparation is classified as dangerous in accordance with Directive 1999/45/EC.

Oxidising

Contact with combustible material may cause fire.

Irritating to respiratory system and skin.

Risk of serious damage to eyes.

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SOLVAY Chemicals





2.2. Potential Health Effects:

Inhalation

- irritation of the upper respiratory tract
- Irritating to mucous membranes
- Repeated or prolonged exposure: Risk of sore throat, nose bleeds.
- (in case of higher concentration): Cough.

Eye contact

- Severe eye irritation
- Lachrymation
- Redness
- Swelling of tissue
- Risk of serious damage to eyes.

- Prolonged skin contact may cause skin irritation.

Ingestion

- Severe irritation
- Irritation of the mouth and throat.
- Symptoms: Nausea, Abdominal pain, Vomiting, Diarrhoea.

Other toxicity effects

- See section 11: Toxicological Information

2.3. Environmental Effects:

- See section 12: Ecological Information

Calcium peroxide

CAS-No. Concentration

1305-79-9 >= 70.0 %

Calcium dihydroxide

CAS-No.

1305-62-0

>= 10.0 - <= 25.0 %

Concentration

Other inorganic calcium compounds Proprietary CAS-No.

>= 10.0 - <= 25.0 % Concentration

4.1. Inhalation

- Remove the subject from dusty environment and let him blow his nose.
- If symptoms persist, call a physician.

- Rinse immediately with plenty of water, also under the eyelids, for at least 15 minutes.
 In the case of difficulty of opening the lids, administer an analgesic eye wash (oxybuprocaine).
- Consult with an ophthalmologist immediately in all cases.

4.3. Skin contact

- Remove and wash contaminated clothing before re-use.
- Wash off with plenty of water.
- If symptoms persist, call a physician.

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2/11



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4.4. Ingestion

- Call a physician immediately.

If victim is conscious:

- If swallowed, rinse mouth with water (only if the person is conscious).
- Do NOT induce vomiting.

If victim is unconscious but breathing:

Artificial respiration and/or oxygen may be necessary.

- 5.1. Suitable extinguishing media
 Use extinguishing measures that are appropriate to local circumstances and the surrounding environment.
 - Wate
 - Water spray

5.2. Extinguishing media which shall not be used for safety reasons

- None

5.3. Special exposure hazards in a fire

- Oxidising
- Oxygen released in thermal decomposition may support combustion
- Contact with combustible material may cause fire.
- Contact with flammables may cause fire or explosions.
- Risk of explosion if heated under confinement.

5.4. Hazardous decomposition products

Oxygen

5.5. Special protective equipment for fire-fighters

- In the event of fire, wear self-contained breathing apparatus.
- Fire fighters must wear fire resistant personnel protective equipment.

5.6. Other information

- Keep product and empty container away from heat and sources of ignition.

6.1. Personal precautions

- Refer to protective measures listed in sections 7 and 8.
- Keep away from incompatible products

6.2. Environmental precautions

- Should not be released into the environment.
- If the product contaminates rivers and lakes or drains inform respective authorities.

6.3. Methods for cleaning up

- Do not add chemical products.
- Pick up and arrange disposal without creating dust.
- All receiving equipment should be clean, vented, dry, labelled and made of material that is compatible with the product.
- Flush with plenty of water.
- Treat recovered material as described in the section "Disposal considerations".

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7.1. Handling

- Clean and dry piping circuits and equipment before any operations.
- Never return unused material to storage receptacle.
- Containers and equipment used to handle the product should be used exclusively for that product.
- Keep away from heat and sources of ignition.
- Keep away from Incompatible products.

7.2. Storage

- Keep in a dry place.
- Keep in a cool, well-ventilated place.
- Keep away from direct sunlight.
- Keep away from heat.
- Keep away from Incompatible products.
- The container must be used exclusively for the product.
- Keep in container fitted with safety valve or vent.

7.3. Packaging material

- Stainless steel
- Plastic material
- glass

7.4. Other information

- Avoid dust formation.
- Refer to protective measures listed in sections 7 and 8.
- In industrial installations, apply the rules for the prevention of major accidents (consult an expert).

Keep away from heat/sparks/open flames/hot surfaces. - No smoking.

8.1. Exposure Limit Values

Calcium peroxide

- SAEL (Solvay Acceptable Exposure Limit) 2007 TWA = 3 mg/m3

- Calcium dihydroxide US. ACGIH Threshold Limit Values 01 2006 time weighted average = 5 mg/m3
- US, OSHA Table Z-1 Limits for Air Contaminants (29 CFR 1910.1000) 02 2006

Permissible exposure limit = 5 mg/m3

Remarks: respirable dust fraction

US. OSHA Table Z-1 Limits for Air Contaminants (29 CFR 1910.1000), 02 2006

Permissible exposure limit = 15 mg/m3

Remarks: Total dust

US. OSHA Table Z-1-A (29 CFR 1910.1000) 1989

time weighted average = 5 mg/m3

US. Tennessee. OELs. Occupational Exposure Limits, Table Z1A 06 2008

time weighted average = 5 mg/m3

Remarks: respirable dust fraction

US. Tennessee. OELs, Occupational Exposure Limits, Table Z1A 06 2008

time weighted average = 15 mg/m3

Remarks: Total dust

Calcium carbonate

- US, OSHA Table Z-1 Limits for Air Contaminants (29 CFR 1910,1000) 02 2006 Permissible exposure limit = 5 mg/m3

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Remarks: respirable dust fraction

US. OSHA Table Z-1 Limits for Air Contaminants (29 CFR 1910.1000) 02 2006

Permissible exposure limit = 15 mg/m3

Remarks: Total dust

- US, OSHA Table Z-1-A (29 CFR 1910.1000) 1989

time weighted average = 5 mg/m3

Remarks: respirable dust fraction

- US. OSHA Table Z-1-A (29 CFR 1910.1000) 1989

time weighted average = 15 mg/m3

Remarks: Total dust

US. Tennessee. OELs. Occupational Exposure Limits, Table Z1A 06 2008

time weighted average = 15 mg/m3

Remarks: Total dust

US. Tennessee. OELs. Occupational Exposure Limits, Table Z1A 06 2008

time weighted average = 5 mg/m3 Remarks: respirable dust fraction

ACGIH® and TLV® are registered trademarks of the American Conference of Governmental Industrial Hygienists.

SAEL = Solvay Acceptable Exposure Limit, Time Weighted Average for 8 hour workdays. No Specific TLV STEL (Short Term Exposure Level) has been set. Excursions in exposure level may exceed 3 times the TLV TWA for no more than a total of 30 minutes during a workday and under no circumstances should they exceed 5 times the TLV TWA.

8.2. Engineering controls

- Ensure adequate ventilation.
- Refer to protective measures listed in sections 7 and 8.
- Apply technical measures to comply with the occupational exposure limits.

8.3. Personal protective equipment

8.3.1. Respiratory protection

- Use only respiratory protection that conforms to international/ national standards.
- Use NIOSH approved respiratory protection.

8.3.2. Hand protection

Wear suitable gloves.

8.3.3. Eye protection

- Chemical resistant goggles must be worn.

8.3.4. Skin and body protection

- Protective suit

8.3.5. Hygiene measures

- Use only in an area equipped with a safety shower.
- Eye wash bottle with pure water
- Handle in accordance with good industrial hygiene and safety practice for diagnostics.

9.1. General Information

Appearance

: Granular

Colour

: light yellow

Odour

: odourless

9.2. Important health safety and environmental information

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: 11.7

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Remarks: saturated aqueous solution

: Remarks: The product is not flammable.

Concentration: 10 g/l Temperature: 20 °C (68 °F)

: Remarks: not applicable

: Remarks: not applicable

: Explosion danger. Remarks: Not explosive : Remarks: Oxidising

Boiling point/boiling range : Remarks: not applicable

Flash point

Flammability

Explosive properties

Oxidizing properties

Vapour pressure

Relative density / Density

Bulk density

Solubility

: 500 kg/m3

: 2.92

: Water 1.65 g/l (calcium hydroxide)

Temperature: 20 °C (68 °F)

: slightly soluble

: Remarks: Decomposes in contact with water.

Partition coefficient: n-octanol/water

: Remarks: not applicable

Vapour density

: Remarks: not applicable

9.3. Other data

Melting point/range

: 275 °C (527 °F)

Remarks: Decomposition

Decomposition temperature

; > 275 °C (527 °F)

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10.1. Stability

- Stable under recommended storage conditions.

10.2. Conditions to avoid

- Exposure to moisture.

- Keep at temperature not exceeding: 275 °C (527 °F)

10.3. Materials to avoid

- Water, Acids, Bases, Heavy metal salts, Reducing agents, Organic materials, Flammable materials

10.4. Hazardous decomposition products

Oxygen

Toxicological data

Acute oral toxicity

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- LD50, rat, > 2,000 mg/kg

Acute inhalation toxicity

LC50, rat, > 5,000 mg/m3

Acute dermal irritation/corrosion

LD50, rat, > 2,000 mg/kg

Skin irritation

- rabbit, No skin irritation

Eye irritation

- Risk of serious damage to eyes.

Sensitisation

- guinea pig, Did not cause sensitization on laboratory animals.

Genetic toxicity in vitro

- In vitro tests did not show mutagenic effects

Remarks

- Risk of serious damage to eyes.

12.1. Ecotoxicity effects

Acute toxicity

- Fishes, Cyprinus carpio, LC50, 48 h, 160 mg/l Crustaceans, Daphnia sp., EC50, 24 h, 25.6 mg/l

12.2. Mobility

<u>Air</u>

Remarks: not applicable

Remarks: low solubility and mobility

Soil/sediments

Remarks: no data available

12.3. Persistence and degradability

Abiotic degradation

Result: not applicable

Water/soil

Result: complexation/precipitation of inorganic materials

Result: non-significant hydrolysis

Biodegradation

- Remarks: The methods for determining biodegradability are not applicable to inorganic substances.

12.4. Bioaccumulative potential

Remarks: not applicable

12.5. Other adverse effects

no data available

- Hazard for the environment is limited due to product properties:
- Aquatic toxicity is unlikely due to low solubility.
- weak solubility and precipitation as carbonate or sulfate in aquatic environment.
- Does not bioaccumulate.

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American International Chemical, Inc. (508) 270-8433 • (800) 238-0001 Framingham, MA 01701

- Diluted product is rapidly neutralized at environmental pH.

13.1. Waste from residues / unused products

- Dilute with plenty of water.
- Dispose of wastes in an approved waste disposal facility.
- Can be landfilled, when in compliance with local regulations.
- In accordance with local and national regulations.

13.2. Packaging treatment

- Clean container with water.
- Empty containers should be taken to an approved waste handling site for recycling or disposal.

- Uncleaned empty packaging
- Dispose of as unused product.
- In accordance with local and national regulations.

13.3. RCRA Hazardous Waste

- Listed RCRA Hazardous Waste (40 CFR 302) No
- Unlisted RCRA Hazardous Waste (40 CFR 302) Yes
- D001 (ignitable waste)

UN-Number	1457
IATA-DGR	
Class	5.1
Packing group	ĬŢ.
ICAO-Labels	Oxidizer
IMDG	
Class	5.1
Packing group	11
ICAO-Labels	Oxidising agent
HI/UN No.	1457
U.S. Dept of Transportation	
Class (Subsidiary) Packing group	5.1 II
Label (Subsidiary)	Oxidising agent
Emergency info:	ERG: 140
Canada (TDG)	
Class (Subsidiary)	5.1
Packing group	
Label (Subsidiary) Emergency info:	Oxidizer ERG: 140
Emergency into.	LING. 140

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15.1. Inventory Information

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Toxic Substance Control Act list (TSCA)	: -		
Australian Inventory of Chemical Substances (AICS)	: -	In compliance with inventory.	
Canadian Domestic Substances List (DSL)	: -	In compliance with inventory.	
Korea Existing Chemicals Inv. (KECI) (KECI (KR))	: -	In compliance with inventory.	
EU list of existing chemical substances (EINECS)		In compliance with inventory.	
Japan (ENCS) List (ENCS (JP))	1 -	In compliance with inventory.	
Inventory of Existing Chemical Substances (China) (IECS)		In compliance with inventory.	
Philippine Inventory of Chemicals and Chemical Substances (PICCS)	1 -	In compliance with inventory.	
New Zealand Inventory (in preparation) (NZ)		In compliance with inventory.	

15.2. Other regulations

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 302 Extremely Hazardous Substance (40 CFR 355, Appendix A)

- not regulated.

SARA Hazard Designation (SARA 311/312)

- Acute Health Hazard: Yes.
- Fire Hazard: Yes.

US. EPA Emergency Planning and Community Right-To-Know Act (EPCRA) SARA Title III Section 313 Toxic Chemicals (40 CFR 372.65) - Supplier Notification Required

- not regulated.

US. EPA CERCLA Hazardous Substances (40 CFR 302)

not regulated

US. New Jersey Worker and Community Right-to-Know Act (New Jersey Statute Annotated Section 34:5A-5)

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US. Pennsylvania Worker and Community Right-to-Know Law (34 Pa. Code Chap. 301-323)

- not regulated.

US. California Safe Drinking Water & Toxic Enforcement Act (Proposition 65)

- not regulated.

15.3. Classification and labelling

Canada. Canadian Environmental Protection Act (CEPA). WHMIS Ingredient Disclosure List (Can. Gaz., Part II, Vol. 122, No. 2)

- C Oxidizing Material
- D2B Toxic Material Causing Other Toxic Effects

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Remarks: This product has been classified in accordance with the hazard criteria of the Controlled Products Regulations and the MSDS contains all the information required by the Controlled Products Regulations.

EC Labe

- The product is classified and labelled in accordance with Directive 1999/45/EC.

Symbol(s)	O Xi	Oxidising Irritant
R-phrase(s)	R 8 R37/38 R41	Contact with combustible material may cause fire. Irritating to respiratory system and skin. Risk of serious damage to eyes.
S-phrase(s)	\$ 3 \$ 8 \$17 \$22 \$24/25 \$26	Keep in a cool place. Keep container dry. Keep away from combustible material. Do not breathe dust. Avoid contact with skin and eyes. In case of contact with eyes, rinse immediately with plenty of water and seek medical advice. Wear suitable gloves and eye/face protection.

Ratings:

NFPA (National Fire Protection Association)

Health = 2 Flammability = 0 Instability = 1 Special = OX

HMIS (Hazardous Material Information System)

Health = 2 Fire = 0 Reactivity = 1 PPE: Supplied by User; dependent on local conditions

Further information

- Update
- This data sheet contains changes from the previous version in section(s): 3, 7.5, 9.3, 10
- Distribute new edition to clients

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